

Since January 2020 Elsevier has created a COVID-19 resource centre with free information in English and Mandarin on the novel coronavirus COVID-19. The COVID-19 resource centre is hosted on Elsevier Connect, the company's public news and information website.

Elsevier hereby grants permission to make all its COVID-19-related research that is available on the COVID-19 resource centre - including this research content - immediately available in PubMed Central and other publicly funded repositories, such as the WHO COVID database with rights for unrestricted research re-use and analyses in any form or by any means with acknowledgement of the original source. These permissions are granted for free by Elsevier for as long as the COVID-19 resource centre remains active. Contents lists available at ScienceDirect





Chaos, Solitons and Fractals Nonlinear Science, and Nonequilibrium and Complex Phenomena

journal homepage: www.elsevier.com/locate/chaos

Stationary distribution and probability density function of a stochastic SVIS epidemic model with standard incidence and vaccination strategies



Baoquan Zhou^a, Daqing Jiang^{a,b,*}, Yucong Dai^a, Tasawar Hayat^{b,c}, Ahmed Alsaedi^b

^a College of Science, China University of Petroleum (East China), Qingdao 266580, P.R. China
^b Nonlinear Analysis and Applied Mathematics(NAAM)-Research Group, Department of Mathematics, King Abdulaziz University, Jeddah, Saudi Arabia

^c Department of Mathematics, Quaid-i-Azam University 45320, Isamabad 44000, Pakistan

ARTICLE INFO

Article history: Received 5 May 2020 Revised 7 November 2020 Accepted 15 December 2020 Available online 24 December 2020

Keywords: Stochastic SVIS epidemic model Vaccination Ergodic stationary distribution Fokker-Planck equation Probability density function Extinction

ABSTRACT

Considering the great effect of vaccination and the unpredictability of environmental variations in nature, a stochastic Susceptible-Vaccinated-Infected-Susceptible (SVIS) epidemic model with standard incidence and vaccination strategies is the focus of the present study. By constructing a series of appropriate Lyapunov functions, the sufficient criterion $\mathscr{R}_0^s > 1$ is obtained for the existence and uniqueness of the ergodic stationary distribution of the model. In epidemiology, the existence of a stationary distribution indicates that the disease will be persistent in a long term. By taking the stochasticity into account, a quasi-endemic equilibrium related to the endemic equilibrium of the deterministic system is defined. By means of the method developed in solving the general three-dimensional Fokker-Planck equation, the exist expression of the probability density function of the stochastic as simplificance, the explicit density function can reflect all dynamical properties of an epidemic system. Next, a simple result of disease extinction is obtained. In addition, several numerical simulations and parameter analyses are performed to illustrate the theoretical results. Finally, the corresponding results and conclusions are discussed at the end of the paper.

© 2020 Elsevier Ltd. All rights reserved.

1. Introduction

1.1. Research background

It is well established that many infectious diseases have a critical influence on global social economies and human health. More precisely, the detailed statistics reported by the World Health Organization (WHO) show that approximately one-third of all deaths worldwide are caused by various epidemics. Recently, the global outbreak of COVID-19 with high transmission has also increased awareness of the importance of preventing and controlling infectious diseases. In epidemiology, mathematical models have provided several effective approaches to describe the characteristics and spread of epidemics in the last hundred years. In 1927, by dividing the population into two clusters, which includes people susceptible to the disease and infected individuals, Kermack and McKendrick [1] initially proposed the classical susceptible-infected-

* Corresponding author. *E-mail address:* daqingjiang2010@hotmail.com (D. Jiang). susceptible (SIS) epidemic model and established the corresponding threshold theory. Since then, various realistic ordinary differential equations (ODEs) have been extended to analyze and control the transmission of diseases [2–8]. For instance, Hove-Musekwa and Nyabadza [4] developed a HIV/AIDS model with active screening of disease carriers and obtained the corresponding basic reproduction number. Considering the effect of vertical infection, Tuncer and Martcheva [6] formulated a hepatitis B model with acute infection and carriers.

With the accelerated development of science and technology, vaccination comprises a common precaution that reduces the infection rate and even immunizes against some contagious diseases, such as measles, cholera, and tuberculosis [9]. According to a 2005 WHO report, the eradication of smallpox has been considered the most spectacular success of routine vaccination. Thus, some basic epidemic models with vaccination strategies have been studied in the last several decades [9–13]. In [9], Liu et al. obtained the global stability of equilibria and analyzed the effect of pulse vaccination. Gao et al. [11] proposed mixed vaccination strategies in the SIRS epidemic model with seasonal variability on infection. However,

the vast majority of infection processes are caused by person-toperson contact. As Ma and Wang [2] described, the classical bilinear incidence rate is reasonably assumed by the simple massaction law. From Anderson and May [14], Hethcote [15], this law is a good approximation for some communicable diseases, such as dengue fever and avian influenza. However, studies (see, e.g., [16,17]) have shown that the underlying assumption of homogeneous mixing and homogeneous environmental for several sexually transmitted diseases, e.g., HIV/AIDS and syphilis, may be invalid. In addition, owing to the psychological effect, susceptible individuals may tend to reduce the number of contacts with the infected per unit time as the numbers of the infected individuals increase [18,19]. As a result, the corresponding adequate incidence rate should be modified as a nonlinear form. More importantly, Anderson and May [20,21] pointed out that various epidemic models with standard incidence are suitable for human beings and some gregarious animals.

Given the above, a SVIS epidemic system with standard incidence and vaccination is the focus of the present study.

1.2. Deterministic SVIS model and dynamical properties

The total population N(t) is divided into three compartments, namely susceptible people S(t), infected individuals I(t), and vaccinees V(t) that are in the vaccination process at time t. Then, the corresponding deterministic SVIS epidemic model with standard incidence and vaccination strategies takes the form

$$\begin{cases} \frac{dS(t)}{dt} = \Lambda - (\mu + \vartheta)S - \frac{\beta SI}{N} + \gamma V + \delta I, \\ \frac{dV(t)}{dt} = \vartheta S - (\mu + \gamma)V, \\ \frac{dI(t)}{dt} = \frac{\beta SI}{N} - (\mu + \alpha + \delta)I, \end{cases}$$
(1.1)

where Λ denotes the recruitment rate of the susceptible, β is the effective contact rate, μ depicts the natural death rate of the population, α denotes the additional death rate due to the disease, ϑ is the vaccination rate of the susceptible, γ denotes the immunity loss coefficient of the vaccinated, and δ reflects the recovery rate of the infected. In epidemiology, these biological parameters are assumed to be positive.

Following similar results described by Ma and Zhou [22], the corresponding basic reproduction number of system (1.1) takes the form

$$\mathscr{R}_{0} = \frac{\beta(\mu + \gamma)}{(\mu + \gamma + \vartheta)(\mu + \alpha + \delta)}.$$
(1.2)

By defining a positive invariant set $\mathcal{D}_0 = \{(S, V, I) | S \ge 0, V \ge 0, I \ge 0, S + V + I \le \frac{\Lambda}{\mu}\}$, two possible equilibria and their dynamical properties are then given as follows.

• Assuming that $\mathscr{R}_0 \leq 1$, the disease-free, $E_0 = (S^0, V^0, I^0) = \left(\frac{\Lambda(\mu+\gamma)}{\mu(\mu+\gamma+\vartheta)}, \frac{\Lambda\vartheta}{\mu(\mu+\gamma+\vartheta)}, 0\right)$, are then globally asymptotically stable in \mathscr{D}_0 .

• If $\mathscr{R}_0 > 1$, there is a unique endemic equilibrium $E^+ = (S^+, V^+, I^+)$, where $I^+ = \frac{\Lambda(\mathscr{R}_0 - 1)}{\mu + (\mu + \alpha)(\mathscr{R}_0 - 1)}$, $S^+ = \frac{\Lambda(\mu + \gamma)}{(\mu + \gamma + \vartheta)[\mu + (\mu + \alpha)(\mathscr{R}_0 - 1)]}$, $V^+ = \frac{\Lambda\vartheta}{(\mu + \gamma + \vartheta)[\mu + (\mu + \alpha)(\mathscr{R}_0 - 1)]}$. Moreover, E^+ is globally asymptotically stable in \mathscr{D}_0 , but E_0 is unstable.

1.3. Stochastic SVIS epidemic model

In fact, Truscott and Gilligan [23] pointed out that the spread of infection, travel of populations, and design of control strategies are critically perturbed by some environmental variations. Therefore, it is more reasonable to construct a corresponding stochas-

tic model to reveal the epidemiological characteristics of infectious diseases by comparison with the deterministic model. Notably, there are various possible approaches to simulate the random effects from biological significance and mathematical perspective [24]. For instance, making use of the fatal properties and multiplex networks, Zhu et al. [25], Jia et al. [26] studied the SIR epidemic spreading process and analyzed individual decisionmaking behavior. In 2002, the most classical assumption that random changes always fluctuate around some average values due to continuous disturbances in nature, adopted by Mao et al. [27], became a common way of describing environmental variations. Moreover, the above random fluctuations are all assumed to be types of white noise. Therefore, many authors have formulated the relevant stochastic differential equations (SDEs) with linear noises for the transmission of various epidemics [28-36]. As an example, Qi and Jiang [29] studied the impact of virus carrier screening and actively seeking treatment on the dynamical behavior of a stochastic HIV/AIDS epidemic model with bilinear incidence. In [34], Shi and Zhang focused on the corresponding stochastic avian influenza system and investigated the existence of the unique ergodic stationary distribution. In addition, several dynamical analyses of the stochastic SIS models or epidemic systems with vaccination have been conducted [37-40]. In [37], Zhao and Jiang creatively proposed a general theory about extinction and persistence in mean based on a stochastic SIS epidemic model with vaccination. Zhang and Jiang [39] obtained sufficient conditions for a stochastic SIS system with saturated incidence and double epidemic diseases. By taking the periodicity effect into account, they still investigated a stochastic SVIR epidemic model with vaccination strategies, and derived the criteria for the existence of non-trivial positive periodic solution [40]. Given the above, in the present study it is assumed that the environmental noises are separately proportional to the compartments S, V and I. Then, the corresponding system (1.1) with the stochastic perturbations is described by

$$\begin{cases} dS(t) = \left[\Lambda - \left(\mu + \vartheta\right)S - \frac{\beta SI}{N} + \gamma V + \delta I\right]dt + \sigma_1 SdB_1(t), \\ dV(t) = \left[\vartheta S - (\mu + \gamma)V\right]dt + \sigma_2 VdB_2(t), \\ dI(t) = \left[\frac{\beta SI}{N} - (\mu + \alpha + \delta)I\right]dt + \sigma_3 IdB_3(t), \end{cases}$$
(1.3)

where $B_1(t)$, $B_2(t)$ and $B_3(t)$ are three independent standard Brownian motions (or Wiener processes), with $\sigma_i^2 > 0$ (i = 1, 2, 3) denoting their intensities.

From the perspective of biomathematics, the existence and ergodicity of stationary distribution indicates that an infectious disease will prevail and persist in long-term development. More importantly, the corresponding probability density function of the stationary distribution can reflect all statistical properties of the individuals *S*, *V* and *I*. It can be regarded as a great intersection of epidemiological dynamics and statistics. It should be pointed out that there are relatively few studies devoted to the explicit expression of probability density function due to the difficulty of solving the corresponding Fokker-Planck equation. To the best of our ability, several general methods of solving the corresponding algebraic equations are developed herein that are equivalent to the Fokker-Planck equation, and the exact expression of density function is derived.

The rest of this paper is organized follows. In Section 2, several mathematical notations and important lemmas for the dynamical analyses of system (1.3) are presented. The sufficient conditions for the existence and uniqueness of the ergodic stationary distribution of system (1.3) are obtained in Section 3. By means of the developed approaches in solving the general three-dimensional Fokker-

Planck equation, the exact expression of the probability density function for the stationary distribution is derived in Section 4. In Section 5, several simple criteria for the disease extinction of system (1.3) are given. In Section 6, several numerical simulations are performed, together with parameter analyses to validate the theoretical results. Finally, the corresponding result discussions and main conclusions are shown, compared with existing articles, at the end of the paper.

2. Preliminaries and necessary lemmas

Throughout the paper, let $\{\Omega, \mathscr{F}, \{\mathscr{F}_t\}_{t\geq 0}, \mathbb{P}\}\$ be a complete probability space with a filtration $\{\mathscr{F}_t\}_{t\geq 0}$ with a filtration $\{\Gamma_t\}_{t\geq 0}$ satisfying the usual conditions (i.e., it is increasing and right continuous, while Γ_0 contains all \mathbb{P} -null sets. Assuming that $A_{m\times n}$ is a real matrix, let A^{τ} be the transpose matrix of A. If m = n, A^{-1} depicts the inverse matrix of A. The reader is referred to Mao [41] for detailed descriptions. For convenience, let \mathbb{R}^n be the n-dimensional Euclidean space, and

$$\mathbb{R}^n_+ = \{(x_1, ..., x_n) | x_k > 0, 1 \le k \le n\},\\ \Xi_n = \left(\frac{1}{n}, n\right) \times \left(\frac{1}{n}, n\right) \times \left(\frac{1}{n}, n\right).$$

Clearly, the values S, V and I that satisfy system (1.3) are required to be positive for the corresponding dynamical behavior. To this end, the existence of uniqueness of the global positive solution of system (1.3) is described by the following Lemma 2.1.

Lemma 2.1. For any initial value $(S(0), V(0), I(0)) \in \mathbb{R}^3_+$, there is then a unique solution (S(t), V(t), I(t)) of the system (1.3) on $t \ge 0$, and the solution will remain in \mathbb{R}^3_+ with probability 1 (a.s.).

The detailed proof is almost the same as those in Zhou et al. [28], and thus it is omitted here. Next, let X(t) be a homogeneous Markov process defined on \mathbb{R}^n that satisfies the following SDE,

$$dX(t) = \phi(X(t))dt + \sum_{k=1}^{l} g_k(X)dB_k(t),$$

where the diffusion matrix $H(x) = (\bar{a}_{ij}(x))$, and $\bar{a}_{ij}(x) = \sum_{k=1}^{l} g_k^{(i)}(x) g_k^{(j)}(x)$. Then, the corresponding ergodicity theory and the existence of stationary distribution are described by the following Lemma 2.2.

Lemma 2.2. (Has'miniskii [42]) The Markov process X(t) has a unique ergodic stationary distribution $\varpi(\cdot)$, if there exists a bounded domain $\mathbb{D}_0 \subset \mathbb{R}^n$ with a regular boundary Γ and the following are true.

 (\mathscr{A}_1) . In the domain \mathbb{D}_0 and some neighborhood thereof, the smallest eigenvalue of the diffusion matrix H(x) is bounded away from zero.

(\mathscr{A}_2). There is a non-negative C^2 -function U(X(t)) such that $\mathscr{L}V(X(t))$ is negative for any $\mathbb{R}^n \setminus \mathbb{D}_0$.

Then, for any $x \in \mathbb{R}^n$ and integral function $\varphi(\cdot)$ with respect to the measure $\overline{\varpi}(\cdot)$, it follows that

$$\mathbb{P}\left\{\lim_{t\to\infty}\frac{1}{t}\int_0^t\varphi(X(t))dt=\int_{\mathbb{R}^n}\varphi(x)\varpi(dx)\right\}=1.$$

By Zhou et al. [28], two important lemmas of solving the special algebraic equations are given as follows.

Lemma 2.3. ([28]) Let Υ_1 be a symmetric matrix, for the threedimensional algebraic equation $G_0^2 + A_0 \Upsilon_1 + \Upsilon_1 A_0^{\tau} = 0$, where $G_0 = diag(1, 0, 0)$,

$$A_0 = \begin{pmatrix} -p_1 & -p_2 & -p_3 \\ 1 & 0 & 0 \\ 0 & 1 & 0 \end{pmatrix}.$$
 (2.1)

If $p_1 > 0$, $p_3 > 0$ and $p_1p_2 - p_3 > 0$, then Υ_1 is positive definite, which follows

$$\Upsilon_1 = \begin{pmatrix} \frac{p_2}{2(p_1p_2-p_3)} & 0 & -\frac{1}{2(p_1p_2-p_3)} \\ 0 & \frac{1}{2(p_1p_2-p_3)} & 0 \\ -\frac{1}{2(p_1p_2-p_3)} & 0 & \frac{p_1}{2p_3(p_1p_2-p_3)} \end{pmatrix}$$

Lemma 2.4. ([28]) Consider the three-dimensional algebraic equation $G_0^2 + B_0 \Upsilon_2 + \Upsilon_2 B_0^{\tau} = 0$, where Υ_2 is a symmetric matrix, $G_0 = diag(1, 0, 0)$,

$$B_0 = \begin{pmatrix} -q_1 & -q_2 & -q_3 \\ 1 & 0 & 0 \\ 0 & 1 & q_{33} \end{pmatrix}.$$
 (2.2)

Assuming that $q_1 > 0$ and $q_2 > 0$, then Υ_2 is positive semi-definite, which takes the form

$$\Upsilon_2 = \begin{pmatrix} \frac{1}{2q_1} & 0 & 0\\ 0 & \frac{1}{2q_1q_2} & 0\\ 0 & 0 & 0 \end{pmatrix}$$

Finally, combining Lemmas 2.3–2.4 and the Routh-Hurwitz criterion [43], two general theories are developed in solving the similar algebraic equations, i.e., Lemmas 2.5 and 2.6.

Lemma 2.5. For the three-dimensional algebraic equation $G_1^2 + C_0 \Upsilon_3 + \Upsilon_3 C_0^{\tau} = 0$, where Υ_3 is a symmetric matrix, $G_1 = diag(a_0, 0, 0) \ (a_0 \neq 0)$

$$C_0 = \begin{pmatrix} c_{11} & c_{12} & c_{13} \\ 0 & c_{22} & c_{23} \\ 0 & c_{32} & c_{33} \end{pmatrix}.$$
 (2.3)

If $c_{11} < 0$, then Υ_3 is a positive semi-definite matrix of the form

$$\Upsilon_3 = \begin{pmatrix} -\frac{a_0^2}{2c_{11}} & 0 & 0\\ 0 & 0 & 0\\ 0 & 0 & 0 \end{pmatrix}.$$

Lemma 2.6. For any real matrices $A = (a_{ij})_{3\times 3}$, $\Pi = diag(\alpha_1^2, \alpha_2^2, \alpha_3^2)$, where $\alpha_i > 0$ (i = 1, 2, 3). Assume that Σ_0 is a symmetric matrix, for the three-dimensional algebraic equation

$$\Pi + A\Sigma_0 + \Sigma_0 A^\tau = 0. \tag{2.4}$$

By defining the characteristic polynomials of A as $\psi_A(\lambda) = \lambda^3 + r_1\lambda^2 + r_2\lambda + r_3$, if A has all negative real part eigenvalues – that is, $r_1 > 0$, $r_3 > 0$, $r_1r_2 - r_3 > 0$ – then Σ_0 is positive definite.

Remark 1. From Zhou et al. [28], A_0 and B_0 are called standard R_1 and R_2 matrices, respectively. Similarly, it is assumed that C_0 is a standard R_3 matrix. In addition, subsection (I) of Appendix A gives the detailed proof of Lemma 2.5. The corresponding proof of Lemma 2.6 and the special form of Σ_0 are shown in subsection (II) of Appendix A.

3. Stationary distribution and ergodicity of system (1.3)

In this section, the focus is on the sufficient conditions for the existence and ergodicity of stationary distribution for system (1.3). Moreover, one must guarantee that the results have no difference from those in the deterministic system (1.1). Define

$$\mathscr{R}_{0}^{s} = \frac{\mu\beta\left(\mu + \gamma + \frac{\sigma_{2}^{2}}{2}\right)}{\left[\left(\mu + \frac{\sigma_{1}^{2}}{2}\right)\left(\mu + \gamma + \frac{\sigma_{2}^{2}}{2}\right) + \vartheta\left(\mu + \frac{\sigma_{2}^{2}}{2}\right)\right]\left(\mu + \alpha + \delta + \frac{\sigma_{3}^{2}}{2}\right)}.$$
(3.1)

Theorem 3.1. Assuming that $\mathscr{R}_0^s > 1$, then the solution (S(t), V(t), I(t)) of system (1.3) with any initial value $(S(0), V(0), I(0)) \in \mathbb{R}^3_+$ is ergodic and has a unique stationary distribution $\overline{\varpi}(\cdot)$.

Proof. By Lemma 2.1, for any initial value $(S(0), V(0), I(0)) \in \mathbb{R}^3_+$, system (1.3) has a unique global positive solution $(S(t), V(t), I(t)) \in \mathbb{R}^3_+$. Then, the proof of Theorem 3.1 is divided into the following two steps: (i) construct a pair of a C^2 -Lyapunov function U(S, V, I) and bounded domain \mathbb{D}_{ϵ} such that $\mathscr{L}U \leq -1$ for any $(S, V, I) \in \mathbb{R}^3_+ \setminus \mathbb{D}_{\epsilon}$, and (ii) validate the condition (\mathscr{A}_1) of Lemma 2.2. **Step 1.** Consider a suitable C^2 -function W(S, V, I) in the form

$$W(S, V, I) = M_0 (S + V + I - a_1 \ln S - a_1 a_2 \ln V - a_3 \ln I) - \ln S - \ln V + (S + V + I),$$

where a_1, a_2, a_3 are all positive constant and are determined in (3.4), and $M_0 = \frac{\Lambda + 2\mu + \gamma + \vartheta + \frac{\sigma_1^2 + \sigma_2^2}{2} + 2}{3\Lambda(\sqrt[3]{\mathcal{R}_0^2} - 1)} > 0$, which indicates

$$-3M_0\Lambda\left(\sqrt[3]{\mathscr{R}_0^{\rm s}}-1\right) + \Lambda + 2\mu + \gamma + \vartheta + \frac{\sigma_1^2 + \sigma_2^2}{2} = -2.$$
(3.2)

Define, for simplicity,

 $W_1 = S + V + I - a_1 \ln S - a_1 a_2 \ln V - a_3 \ln I, W_2 = -\ln S - \ln V, W_3 = S + V + I.$

Applying Itô's formula to W_1 , which is shown in subsection (III) of Appendix B, obtains

$$\begin{aligned} \mathscr{L}W_{1} &= \Lambda - \mu N - \alpha I - a_{1} \left[\frac{\Lambda}{S} - \frac{\beta I}{N} + \frac{\gamma V}{S} + \frac{\delta I}{S} - \left(\mu + \vartheta + \frac{\sigma_{1}^{2}}{2} \right) \right] \\ &- a_{1}a_{2} \left[\frac{\vartheta S}{V} - \left(\mu + \gamma + \frac{\sigma_{2}^{2}}{2} \right) \right] - a_{3} \left[\frac{\beta S}{N} - \left(\mu + \alpha + \delta + \frac{\sigma_{1}^{2}}{2} \right) \right] \\ &\leq \Lambda - \left(\mu N + \frac{a_{1}\Lambda}{S} + \frac{a_{3}\beta S}{N} \right) + a_{1} \left(\mu + \vartheta + \frac{\sigma_{1}^{2}}{2} \right) + a_{3} \left(\mu + \alpha + \delta + \frac{\sigma_{2}^{2}}{2} \right) \\ &- \left(\frac{a_{1}\gamma V}{S} + \frac{a_{1}a_{2}\vartheta S}{V} \right) + a_{1}a_{2} \left(\mu + \gamma + \frac{\sigma_{2}^{2}}{2} \right) + \frac{a_{1}\beta I}{N} \\ &\leq \Lambda - 3\sqrt[3]{a_{1}a_{3}\Lambda\mu\beta} - 2\sqrt{a_{1}^{2}a_{2}\vartheta\gamma} + a_{1} \left(\mu + \vartheta + \frac{\sigma_{1}^{2}}{2} \right) + a_{3} \left(\mu + \alpha + \delta + \frac{\sigma_{2}^{2}}{2} \right) + a_{1}a_{2} \left(\mu + \gamma + \frac{\sigma_{2}^{2}}{2} \right) + \frac{a_{1}\beta I}{N}. \end{aligned}$$

$$(3.3)$$

Choosing a_1, a_2 and a_3 such that

$$a_2\left(\mu+\gamma+\frac{\sigma_2^2}{2}\right)^2 = \gamma\vartheta, \quad a_1\left[\left(\mu+\vartheta+\frac{\sigma_1^2}{2}\right)-\frac{\gamma\vartheta}{\mu+\gamma+\frac{\sigma_2^2}{2}}\right] = a_3\left(\mu+\alpha+\delta+\frac{\sigma_3^2}{2}\right) = \Lambda, \tag{3.4}$$

which means $a_1 = \frac{\gamma \vartheta}{(\mu + \gamma + \frac{\sigma_2^2}{2})^2}$, $a_2 = \frac{\Lambda}{\mu + \vartheta + \frac{\sigma_1^2}{2} - \frac{\gamma \vartheta}{\mu + \gamma + \frac{\sigma_2^2}{2}}}$ and $a_3 = \frac{\Lambda}{\mu + \alpha + \delta + \frac{\sigma_3^2}{2}}$, one can obtain

$$\mathscr{L}W_{1} \leq 2\Lambda + a_{1} \left[\left(\mu + \vartheta + \frac{\sigma_{1}^{2}}{2} \right) - \frac{\gamma \vartheta}{\mu + \gamma + \frac{\sigma_{2}^{2}}{2}} \right] - 3\sqrt[3]{\frac{a_{1}\Lambda^{2}\mu\beta}{\mu + \alpha + \delta + \frac{\sigma_{2}^{2}}{2}}} + \frac{a_{1}\beta I}{N}$$
$$= 3\Lambda - 3\Lambda\sqrt[3]{\mathscr{R}_{0}^{5}} + \frac{a_{1}\beta I}{N} = -3\Lambda\left(\sqrt[3]{\mathscr{R}_{0}^{5}} - 1\right) + \frac{a_{1}\beta I}{N}.$$
(3.5)

Applying Itô's formula to W_2, W_3 similarly obtains

$$\mathscr{L}W_{2} = \left[-\frac{\Lambda}{S} + \left(\mu + \vartheta + \frac{\sigma_{1}^{2}}{2}\right) + \frac{\beta I}{N} - \frac{\gamma V}{S} - \frac{\delta I}{S}\right] - \frac{\vartheta S}{V} + \left(\mu + \gamma + \frac{\sigma_{2}^{2}}{2}\right)$$

$$\leq -\frac{\Lambda}{S} - \frac{\vartheta S}{V} + \frac{\beta I}{N} + 2\mu + \gamma + \vartheta + \frac{\sigma_{1}^{2} + \sigma_{2}^{2}}{2},$$
(3.6)

$$\mathscr{L}W_3 = \Lambda - \mu(S+I) - (\mu + \alpha)I \le \Lambda - \mu N.$$
(3.7)

Additionally, note that W(S, V, I) is a continuous function satisfying

$$\liminf_{l \to +\infty, (S,V,I) \in \mathbb{R}^3_+ \setminus \Xi_l} W(S,V,I) = +\infty$$

Hence, W(S, V, I) has a minimum value W_0 . Defining a non-negative C^2 -function U(S, V, I): $\mathbb{R}^3_+ \to \mathbb{R}^1_+$ by

$$U(S, V, I) = W(S, V, I) - W_0,$$

and combining (3.2) and (3.5)–(3.7), it can be shown that

$$\mathscr{L}U \leq -3M_0\Lambda\left(\sqrt[3]{\mathscr{R}_0^S} - 1\right) + \frac{a_1M_0\beta I}{N} - \frac{\Lambda}{S} - \frac{\vartheta S}{V} + \frac{\beta I}{N} + \left(\Lambda + 2\mu + \gamma + \vartheta + \frac{\sigma_1^2 + \sigma_2^2}{2}\right) - \mu N$$
$$= -2 + \frac{(a_1M_0 + 1)\beta I}{S + V + I} - \frac{\Lambda}{S} - \frac{\vartheta S}{V} - \mu (S + V + I).$$
(3.8)

Next, the corresponding compact subset \mathbb{D}_{ϵ} is contructed by

$$D_{\epsilon} = \left\{ (S, V, I) \in \mathbb{R}^3_+ \middle| S \ge \epsilon, V \ge \epsilon^2, I \ge \epsilon^3, S + V + I \le \frac{1}{\epsilon} \right\},\$$

where $\epsilon > 0$ is a sufficiently small constant such that the following inequalities hold:

$$-2 + (a_1 M_0 + 1)\beta - \frac{\min(\Lambda, \vartheta, \mu)}{\epsilon} \le -1,$$
(3.9)

$$-2 + (a_1 M_0 + 1)\beta\epsilon \le -1. \tag{3.10}$$

For convenience, consider the following four subsets of $\mathbb{R}^3_+ \setminus \mathbb{D}_{\epsilon}$:

$$\mathbb{D}_{1,\epsilon}^{c} = \left\{ (S, V, I) \in \mathbb{R}_{+}^{3} \middle| S < \epsilon \right\}, \quad \mathbb{D}_{2,\epsilon}^{c} = \left\{ (S, V, I) \in \mathbb{R}_{+}^{3} \middle| V < \epsilon^{2}, S \ge \epsilon \right\},$$

$$\mathbb{D}_{3,\epsilon}^{c} = \left\{ (S, V, I) \in \mathbb{R}_{+}^{3} \middle| I < \epsilon^{3}, V \ge \epsilon^{2} \right\}, \quad \mathbb{D}_{4,\epsilon}^{c} = \left\{ (S, V, I) \in \mathbb{R}_{+}^{3} \middle| S + V + I > \frac{1}{\epsilon} \right\}$$

Now, it must be shown that

 $\mathscr{L}U \leq -1, \quad \forall \ (S, V, I) \in \mathbb{D}_{i,\epsilon} \ (i = 1, 2, 3, 4).$ Case 1. If $(S, V, I) \in \mathbb{D}_{1,\epsilon}$, by (3.8)–(3.9), one can derive

$$\mathscr{L}U \leq -2 + (a_1M_0 + 1)\beta - \frac{\Lambda}{S} \leq -2 + (a_1M_0 + 1)\beta - \frac{\min(\Lambda, \vartheta, \mu)}{\epsilon} \leq -1.$$

Case 2. For any $(S, V, I) \in \mathbb{D}_{2,\epsilon}$, it follows from (3.8)–(3.9) that

$$\mathscr{L}U \leq -2 + (a_1M_0 + 1)\beta - \frac{\vartheta S}{V} \leq -2 + (a_1M_0 + 1)\beta - \frac{\min(\Lambda, \vartheta, \mu)}{\epsilon} \leq -1$$

Case 3. Assuming that $(S, V, I) \in \mathbb{D}_{3,\epsilon}$, by (3.8)–(3.9) it can be seen that

$$\mathscr{L}U \leq -2 + \frac{(a_1M_0+1)\beta I}{V} \leq -2 + (a_1M_0+1)\beta\epsilon \leq -1.$$

Case 4. If $(S, V, I) \in \mathbb{D}_{1,\epsilon}$, from (3.8) and (3.10), one has

$$\mathscr{L}U \leq -2 + (a_1M_0 + 1)\beta - \mu(S + V + I) \leq -2 + (a_1M_0 + 1)\beta - \frac{\min(\Lambda, \vartheta, \mu)}{\epsilon} \leq -1$$

Notably, $\mathbb{R}^3_+ \setminus \mathbb{D}_{\epsilon} = \bigcup_{i=1}^4 \mathbb{D}_{i,\epsilon}$. Hence, one equivalently obtains

$$\mathscr{L}U \leq -1, \quad \forall \ (S, V, I) \in \mathbb{R}^3_+ \setminus \mathbb{D}_{\epsilon}.$$

Therefore, the condition (\mathscr{A}_2) of Lemma 2.2 holds. The corresponding diffusion matrix is given by

$$H = \begin{pmatrix} \sigma_1^2 S^2 & 0 & 0 \\ 0 & \sigma_2^2 V^2 & 0 \\ 0 & 0 & \sigma_3^2 l^2 \end{pmatrix}.$$

Clearly, *H* is a positive-definite matrix. Then, the assumption (\mathscr{A}_1) of Lemma 2.2 also holds.

Given the above, the global positive solution (S(t), V(t), I(t)) of system (1.3) follows a unique ergodic stationary distribution $\varpi(\cdot)$. This completes the proof of Theorem 3.1. \Box

Remark 2. Under $\mathscr{R}_0^s > 1$, the existence of the ergodic stationary distribution for system (1.3) is derived. This reveals that the contagious disease will prevail and persist in a population. Furthermore, from the expressions of \mathscr{R}_0^s and \mathscr{R}_0 , it can be obtained that $\mathscr{R}_0^s \leq \mathscr{R}_0$, and the sign holds if and only if $\sigma_1 = \sigma_2 = \sigma_3 = 0$. Consequently, not only does this reveal that random fluctuations have a critical effect on the spread of epidemic, but it also indicates that \mathscr{R}_0^s is a unified threshold of the disease persistence of systems (1.1) and (1.3).

4. Probability density function analysis

By Theorem 3.1, one obtains that the global solution (S(t), V(t), I(t)) of system (1.3) follows a unique ergodic stationary distribution $\varpi(\cdot)$. This section is devoted to deriving the explicit expression of the density function of the distribution $\varpi(\cdot)$ while $\mathscr{R}_0^s > 1$. In fact, the result will present a wide range of possibilities for the further development of epidemiological dynamics. Before this, two necessary transformations of system (1.3) should be first introduced.

4.1. Two important transformations of system (1.3)

(I) (Logarithmic transformation): Let $(u_1, u_2, u_3)^{\tau} = (\ln S, \ln V, \ln I)^{\tau}$, Employing Itô's formula, it follows from system (1.3) that

$$\begin{cases} du_{1} = \left[\Lambda e^{-u_{1}} - \left(\mu + \vartheta + \frac{\sigma_{1}^{2}}{2}\right) - \frac{\beta e^{u_{3}}}{e^{u_{1}} + e^{u_{2}} + e^{u_{3}}} + \gamma e^{u_{2}-u_{1}} + \delta e^{u_{3}-u_{1}}\right] dt + \sigma_{1} dB_{1}(t), \\ du_{2} = \left[\vartheta e^{u_{1}-u_{2}} - \left(\mu + \gamma + \frac{\sigma_{2}^{2}}{2}\right)\right] dt + \sigma_{2} dB_{2}(t), \\ du_{3} = \left[\frac{\beta e^{u_{1}}}{e^{u_{1}} + e^{u_{2}} + e^{u_{3}}} - \left(\mu + \alpha + \delta + \frac{\sigma_{3}^{2}}{2}\right)\right] dt + \sigma_{3} dB_{3}(t). \end{cases}$$
(4.1)

(4.4)

By taking the random effect into consideration, another critical value is defined: $\mathscr{R}_0^c = \frac{\beta(\mu+\gamma+\frac{\sigma_2^2}{2})}{(\mu+\gamma+\epsilon+\frac{\sigma_2^2}{2})(\mu+\alpha+\delta+\frac{\sigma_3^2}{2})}$. Moreover, if $\mathscr{R}_0^c > 1$, then the quasi-stable equilibrium $E^* = (S^*, V^*, I^*) := (e^{u_1^*}, e^{u_2^*}, e^{u_3^*}) \in \mathbb{R}_+^3$ is determined by the following Eq. (4.2):

$$\begin{cases} \Lambda e^{-u_{1}^{*}} - \left(\mu + \vartheta + \frac{\sigma_{1}^{2}}{2}\right) - \frac{\beta e^{u_{3}^{*}}}{e^{u_{1}^{*}} + e^{u_{2}^{*}} + e^{u_{3}^{*}}} + \gamma e^{u_{2}^{*} - u_{1}^{*}} + \delta e^{u_{3}^{*} - u_{1}^{*}} = 0, \\ \vartheta e^{u_{1}^{*} - u_{2}^{*}} - \left(\mu + \gamma + \frac{\sigma_{2}^{2}}{2}\right) = 0, \\ \frac{\beta e^{u_{1}^{*}}}{e^{u_{1}^{*}} + e^{u_{3}^{*}}} - \left(\mu + \alpha + \delta + \frac{\sigma_{3}^{2}}{2}\right) = 0. \end{cases}$$
(4.2)

For convenience, let $\mu_i = \mu + \frac{\sigma_i^2}{2}$ for any i = 1, 2, 3. As a result, it can be derived by (4.2) that $V^* = \frac{\vartheta S^*}{\mu_2 + \gamma}$, $I^* = \frac{(\mu_2 + \gamma + \vartheta)(\mathscr{R}_0^c - 1)S^*}{\mu_2 + \gamma}$, $N^* = S^* + V^* + I^* = \frac{\beta S^*}{\mu_3 + \alpha + \delta}$, where $S^* = \frac{\Lambda(\mu_2 + \gamma)}{\mu_1(\mu_2 + \gamma) + \vartheta \mu_2 + (\mu_3 + \alpha)(\mu_2 + \gamma + \vartheta)(\mathscr{R}_0^c - 1)}$.

Notably, it is easily obtained that $\mathscr{R}_0^s < \mathscr{R}_0^c$. This then indicates that $E^* \in \mathbb{R}^3_+$ if $\mathscr{R}_0^s > 1$. In addition, if there is no stochastic noise in system (1.3), i.e., model (1.1), then $E^* = E^+ = (S^+, V^+, I^+)$.

(II) (Equilibrium offset transformation): Given the above, let $X = (x_1, x_2, x_3)^{\tau} = (u_1 - u_1^*, u_2 - u_2^*, u_3 - u_3^*)$; thus, the corresponding linearized system of (4.1) takes the form

$$\begin{cases} dx_1 = (-a_{11}x_1 + a_{12}x_2 + a_{13}x_3)dt + \sigma_1 dB_1(t), \\ dx_2 = (a_{21}x_1 - a_{21}x_2)dt + \sigma_2 dB_2(t), \\ dx_3 = [(a_{32} + a_{33})x_1 - a_{32}x_2 - a_{33}x_3]dt + \sigma_3 dB_3(t), \end{cases}$$

$$(4.3)$$

where

$$\begin{aligned} a_{11} &= \frac{\Lambda + \gamma V^* + \delta I^*}{S^*} - \frac{\beta S^* I^*}{(N^*)^2}, \quad a_{12} &= \frac{\gamma V^*}{S^*} + \frac{\beta V^* I^*}{(N^*)^2} > 0, \quad a_{13} &= \frac{\delta I^*}{S^*} - \frac{\beta (S^* + V^*) I^*}{(N^*)^2} \\ a_{21} &= \mu_2 + \gamma > 0, \quad a_{32} &= \frac{\beta S^* V^*}{(N^*)^2} > 0, \quad a_{33} &= \frac{\beta S^* I^*}{(N^*)^2} > 0. \end{aligned}$$

4.2. Density function of stationary distribution $\overline{\varpi}(\cdot)$

Theorem 4.1. For any initial value $(S(0), V(0), I(0)) \in \mathbb{R}^3_+$, if $\mathscr{R}^s_0 > 1$, then the stationary distribution $\varpi(\cdot)$ around E^* follows a unique log-normal probability density function $\Phi(S, V, I)$, which is given by

$$\Phi(S,V,I) = (2\pi)^{-\frac{3}{2}} |\Sigma|^{-\frac{1}{2}} e^{-\frac{1}{2}(\ln\frac{S}{S^*},\ln\frac{V}{V^*},\ln\frac{I}{V^*})\Sigma^{-1}(\ln\frac{S}{S^*},\ln\frac{V}{V^*},\ln\frac{I}{V^*})^{\mathrm{r}}},$$

where Σ is a positive definite matrix, and the special form of Σ is given as follows. (1). If $m_1 \neq 0$, $m_2 \neq 0$ and $a_{13} \neq 0$, then $\Sigma = \rho_1^2 (H_1 J_1)^{-1} \Theta_0 [(H_1 J_1)^{-1}]^{\tau} + \rho_2^2 (H_2 J_3 J_2)^{-1} \Theta_0 [(H_2 J_3 J_2)^{-1}]^{\tau} + \rho_3^2 (H_3 J_4)^{-1} \Theta_0 [(M_3 J_3)^{-1}]^{\tau}.$ (2). If $m_1 \neq 0$, $m_2 \neq 0$ and $a_{13} = 0$, then $\Sigma = \varrho_1^2 (H_1 J_1)^{-1} \Theta_0 [(H_1 J_1)^{-1}]^{\tau} + \varrho_2^2 (H_2 J_3 J_2)^{-1} \Theta_0 [(H_2 J_3 J_2)^{-1}]^{\tau} + J_4^{-1} \Theta_4 (J_4^{-1})^{\tau}.$ (3). If $m_1 \neq 0$, $m_2 = 0$ and $a_{13} \neq 0$, then $\Sigma = \rho_1^2 (H_1 J_1)^{-1} \Theta_0 [(H_1 J_1)^{-1}]^{\tau} + a_{32}^2 \sigma_2^2 (\widetilde{H}_2 J_3 J_2)^{-1} \Theta_3 [(\widetilde{H}_2 J_3 J_2)^{-1}]^{\tau} + \rho_3^2 (H_3 J_4)^{-1} \Theta_0 [(M_3 J_3)^{-1}]^{\tau}.$ (4). If $m_1 \neq 0$ and $m_2 = a_{13} = 0$, then $\Sigma = \rho_1^2 (H_1 J_1)^{-1} \Theta_0 [(H_1 J_1)^{-1}]^{\tau} + a_{32}^2 \sigma_2^2 (\widetilde{H}_2 J_3 J_2)^{-1} \Theta_3 [(\widetilde{H}_2 J_3 J_2)^{-1}]^{\tau} + J_4^{-1} \Theta_4 (J_4^{-1})^{\tau}.$ (5). If $m_1 = 0$, $m_2 \neq 0$ and $a_{13} \neq 0$, then $\Sigma = a_{21}^2 \sigma_1^2 (\widetilde{H}_1 J_1)^{-1} \Theta_1 [(\widetilde{H}_1 J_1)^{-1}]^{\tau} + \rho_2^2 (H_2 J_3 J_2)^{-1} \Theta_0 [(H_2 J_3 J_2)^{-1}]^{\tau} + \rho_3^2 (H_3 J_4)^{-1} \Theta_0 [(M_3 J_3)^{-1}]^{\tau}.$ (6). If $m_1 = a_{13} = 0$ and $m_2 \neq 0$, then $\Sigma = a_{21}^2 \sigma_1^2 (\widetilde{H}_1 J_1)^{-1} \Theta_1 [(\widetilde{H}_1 J_1)^{-1}]^{\tau} + \rho_2^2 (H_2 J_3 J_2)^{-1} \Theta_0 [(H_2 J_3 J_2)^{-1}]^{\tau} + J_4^{-1} \Theta_4 (J_4^{-1})^{\tau}.$ (7). If $m_1 = m_2 = 0$ and $a_{13} \neq 0$, then $\Sigma = a_{21}^2 \sigma_1^2 (\widetilde{H}_1 J_1)^{-1} \Theta_1 [(\widetilde{H}_1 J_1)^{-1}]^{\tau} + a_{32}^2 \sigma_2^2 (\widetilde{H}_2 J_3 J_2)^{-1} \Theta_3 [(\widetilde{H}_2 J_3 J_2)^{-1}]^{\tau} + \varrho_3^2 (H_3 J_4)^{-1} \Theta_0 [(M_3 J_3)^{-1}]^{\tau}.$ (8). If $m_1 = m_2 = a_{13} = 0$, then $\Sigma = a_{21}^2 \sigma_1^2 (\widetilde{H}_1 J_1)^{-1} \mathcal{O}_1 [(\widetilde{H}_1 J_1)^{-1}]^\tau + a_{32}^2 \sigma_2^2 (\widetilde{H}_2 J_3 J_2)^{-1} \mathcal{O}_3 [(\widetilde{H}_2 J_3 J_2)^{-1}]^\tau + J_4^{-1} \mathcal{O}_4 (J_4^{-1})^\tau,$ $J_1 = \begin{pmatrix} 1 & 0 & 0 \\ 0 & 1 & 0 \\ 0 & -\frac{a_{32}+a_{33}}{a_{22}} & 1 \end{pmatrix}, \quad J_2 = \begin{pmatrix} 0 & 1 & 0 \\ 0 & 0 & 1 \\ 1 & 0 & 0 \end{pmatrix}, \quad J_3 = \begin{pmatrix} 1 & 0 & 0 \\ 0 & 1 & 0 \\ 0 & \frac{a_{12}}{a_{32}} & 1 \end{pmatrix}, \quad J_4 = \begin{pmatrix} 0 & 0 & 1 \\ 1 & 0 & 0 \\ 0 & 1 & 0 \end{pmatrix},$

$$\begin{split} & \Theta_{0} = \begin{pmatrix} \frac{r_{2}}{2(r_{1}r_{2}-r_{3})} & 0 & -\frac{1}{2(r_{1}r_{2}-r_{3})} \\ 0 & \frac{1}{2(r_{1}r_{2}-r_{3})} & 0 \\ -\frac{1}{2(r_{1}r_{2}-r_{3})} & 0 & \frac{r_{1}}{2r_{3}(r_{1}r_{2}-r_{3})} \end{pmatrix}, \quad \Theta_{1} = \begin{pmatrix} \frac{1}{2w_{1}} & 0 & 0 \\ 0 & \frac{1}{2w_{1}w_{2}} & 0 \\ 0 & 0 & 0 \end{pmatrix}, \quad \Theta_{3} = \begin{pmatrix} \frac{1}{2w_{3}} & 0 & 0 \\ 0 & \frac{1}{2w_{3}w_{4}} & 0 \\ 0 & 0 & 0 \end{pmatrix} \\ & H_{1} = \begin{pmatrix} a_{21}m_{1} & -(a_{21}+a_{33})m_{1} & a_{33}^{2} \\ 0 & m_{1} & -a_{33} \\ 0 & 0 & 1 \end{pmatrix}, \quad \widetilde{H}_{2} = \begin{pmatrix} -a_{32} & -a_{33}-a_{12} - \frac{a_{12}a_{33}}{a_{32}} & a_{32}+a_{33} \\ 0 & 1 & 0 \\ 0 & 0 & 1 \end{pmatrix}, \\ & H_{2} = \begin{pmatrix} -a_{32}m_{2} & -(a_{11}+a_{33})m_{2} & (a_{11}-a_{12} - \frac{a_{12}a_{33}}{a_{32}})^{2} + m_{2}(a_{32}+a_{33}) \\ 0 & 0 & 1 \end{pmatrix}, \quad \Theta_{4} = \begin{pmatrix} \frac{1}{2a_{33}} & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{pmatrix}, \\ & \widetilde{H}_{1} = \begin{pmatrix} a_{21} & -a_{21} & 0 \\ 0 & 1 & 0 \\ 0 & 0 & 1 \end{pmatrix}, \quad H_{3} = \begin{pmatrix} a_{13}a_{21} & -a_{21}(a_{11}+a_{21}) & a_{21}(a_{12}+a_{21}) \\ 0 & 0 & 1 \end{pmatrix}, \end{split}$$

and

$$m_1 = \frac{a_{33}(a_{21} - a_{32} - a_{33})}{a_{21}}, \quad m_2 = a_{13} + \frac{a_{12}(a_{11} - a_{33})}{a_{32}} - \frac{a_{12}^2(a_{32} + a_{33})}{a_{32}^2},$$

 $\varrho_1 = a_{21}m_1\sigma_1, \quad \varrho_2 = -a_{32}m_2\sigma_2, \quad \varrho_3 = a_{13}a_{21}\sigma_3,$

 $r_1 = a_{11} + a_{21} + a_{33}, \quad r_2 = a_{21}(a_{11} - a_{12} + a_{33}) + [a_{11}a_{33} - a_{13}(a_{32} + a_{33})], \quad r_3 = a_{21}a_{33}(a_{11} - a_{12} - a_{13}),$

$$w_1 = a_{11} + a_{21}, \quad w_2 = a_{21}(a_{11} - a_{12} - a_{13}), \quad w_3 = a_{12} + a_{21} + a_{33} + \frac{a_{12}a_{33}}{a_{32}}, \quad w_4 = \frac{a_{21}a_{32}a_{33}(a_{11} - a_{12} - a_{13})}{a_{32}(a_{11} - a_{12}) - a_{12}a_{33}}$$

Proof. For convenience and simplicity, let $B(t) = (B_1(t), B_2(t), B_3(t))^{\tau}$ and

$$M = \begin{pmatrix} \sigma_1^2 & 0 & 0 \\ 0 & \sigma_2^2 & 0 \\ 0 & 0 & \sigma_3^2 \end{pmatrix}, \quad A = \begin{pmatrix} -a_{11} & a_{12} & a_{13} \\ a_{21} & -a_{21} & 0 \\ a_{32} + a_{33} & -a_{32} & -a_{33} \end{pmatrix}.$$

Hence, system (4.3) can be rewritten as dX = AXdt + MdB(t). By the theory of Gardiner [44], the unique density function $\Phi(X)$ around the quasi-endemic equilibrium E^* satisfies the following Fokker-Plauck equation:

$$-\sum_{k=1}^{3} \frac{\sigma_{k}^{2}}{2} \frac{\partial^{2} \Phi}{\partial x_{k}^{2}} + \frac{\partial}{\partial x_{1}} \Big[(-a_{11}x_{1} + a_{12}x_{2} + a_{13}x_{3})\Phi \Big] + \frac{\partial}{\partial x_{2}} \Big[(a_{21}x_{1} - a_{21}x_{2})\Phi \Big] \\ + \frac{\partial}{\partial x_{3}} \Big[((a_{32} + a_{33})x_{1} - a_{32}x_{2} - a_{33}x_{3})\Phi \Big] = 0.$$

$$(4.5)$$

Since the diffusion matrix *M* is a constant matrix, Roozen [45] pointed out that $\Phi(X)$ can be described by a quasi-Gaussian distribution, i.e., $\Phi(X) = c_0 e^{-\frac{1}{2}XQX^{\tau}}$, where $c_0 > 0$ is determined by the normalized condition $\int_{R^3_+} \Phi(X) dX = 1$ and *Q* is a symmetric matrix.

Substituting these results into (4.5), one can obtain that Q obeys the algebraic equation $QM^2Q + A^{\tau}Q + QA = 0$. If Q is a inverse matrix, by letting $\Sigma = Q^{-1}$, an equivalent equation is given by

$$M^2 + A\Sigma + \Sigma A^{\tau} = 0. \tag{4.6}$$

Next, it will be proved that *A* has all negative real-part eigenvalues. The characteristic polynomial of *A* is defined as $\psi(\lambda) = \lambda^3 + p_1\lambda^2 + p_2\lambda + p_3$, where

 $r_1 = a_{11} + a_{21} + a_{33}, \quad r_2 = a_{21}(a_{11} - a_{12} + a_{33}) + [a_{11}a_{33} - a_{13}(a_{32} + a_{33})], \quad r_3 = a_{21}a_{33}(a_{11} - a_{12} - a_{13}).$

By the expressions of S^* , V^* , I^* and N^* , it can be shown that

$$\begin{array}{ll} \text{(i).} & a_{11} = (\mu_1 + \vartheta) + \frac{\beta I^*}{N^*} - \frac{\beta S^* I^*}{(N^*)^2} = (\mu_1 + \vartheta) + \frac{\beta (V^* + I^*) I^*}{(N^*)^2} > 0, \\ \text{(ii).} & a_{11} - a_{12} - a_{13} = \left[\frac{\Lambda + \gamma V^* + \delta I^*}{S^*} - \frac{\beta S^* I^*}{(N^*)^2} \right] - \left[\frac{\gamma V^*}{S^*} + \frac{\beta V^* I^*}{(N^*)^2} \right] - \left[\frac{\delta I^*}{S^*} - \frac{\beta (S^* + V^*) I^*}{(N^*)^2} \right] = \frac{\Lambda}{S^*} > 0, \\ \text{(iii).} & a_{12}a_{33} - a_{13}a_{32} = \frac{\beta S^*}{(N^*)^2} \left(\frac{\gamma V^* I^*}{S^*} + \frac{\beta V^* I^*}{N^*} - \frac{\delta V^* I^*}{S^*} \right) = \frac{(\mu_3 + \gamma + \alpha)\beta V^* I^*}{(N^*)^2} > 0, \\ \end{array}$$

(iv).
$$a_{11} - a_{12} + a_{33} > \frac{\Lambda}{S^*} + \frac{\delta I^*}{S^*} - \frac{\beta V^* I^*}{(N^*)^2} > \frac{(\mu_3 + \alpha + \delta)(\mathscr{R}_0^c - 1)}{\mu_2 + \gamma} \left[(\mu_2 + \gamma) + \frac{\vartheta(\mathscr{R}_0^c - 1)}{\mathscr{R}_0^c} \right] > 0.$$

Consequently, it follows from (i)-(iv) that

$$\begin{array}{ll} (1). & r_1 = a_{11} + a_{21} + a_{33} > 0, & r_3 = a_{21}a_{33}(a_{11} - a_{12} - a_{13}) > 0, \\ (2). & r_2 = a_{21}(a_{11} - a_{12} + a_{33}) + [a_{11}a_{33} - a_{13}(a_{32} + a_{33})] \\ & > (a_{12} + a_{13})a_{33} - a_{13}(a_{32} + a_{33}) \\ & = a_{12}a_{33} - a_{13}a_{32} > 0. \end{array}$$

(4.7)

$$\begin{array}{ll} (3). & r_1r_2 - r_3 = (a_{11} + a_{21} + a_{33})\{a_{21}(a_{11} - a_{12} + a_{33}) + [a_{11}a_{33} - a_{13}(a_{32} + a_{33})]\} - a_{21}a_{33}(a_{11} - a_{12} - a_{13}) \\ & = a_{11}r_2 + a_{21}[(a_{11} + a_{33})a_{33} - a_{13}a_{32}] + a_{33}[a_{11}a_{33} - a_{13}(a_{32} + a_{33})] \\ & = a_{11}r_2 + a_{21}a_{33}(a_{11} - a_{12} + a_{33}) + a_{33}^2(a_{11} - a_{12} - a_{13}) + (a_{21} + a_{33})(a_{12}a_{33} - a_{13}a_{32}) \\ & > a_{11}r_2 > 0. \end{array}$$

Combining the above (1)-(3) and Lemma 2.6, that Σ of Eq. (4.6) is positive definite can be derived.

However, following the corresponding proof of Lemma 2.6, which is shown in subsection (II) of Appendix A, the exact expression of Σ is given. First, by the finite independent superposition principle, (4.6) can be equivalently transformed into the sum of solution to the following algebraic sub-equations,

$$M_k^2 + A\Sigma_k + \Sigma_k A^\tau = 0,$$

where $M_1 = diag(\sigma_1, 0, 0)$, $M_2 = diag(0, \sigma_2, 0)$, $M_3 = diag(0, 0, \sigma_3)$, and the symmetric matrices Σ_k (k = 1, 2, 3) are their respective solutions of the symmetric matrices Σ_k (k = 1, 2, 3) are the symmetric matrices Σ_k (k = 1, 2, 3). tions. Clearly, $\Sigma = \Sigma_1 + \Sigma_2 + \Sigma_3$. Now, the special expression of Σ are derived by the following three steps. Step 1. For the algebraic equation

$$M_1^2 + A\Sigma_1 + \Sigma_1 A^\tau = 0,$$

denote $A_1 = J_1 A J_1^{-1}$, where the elimination matrix J_1 and A_1 are derived by

$$J_1 = \begin{pmatrix} 1 & 0 & 0 \\ 0 & 1 & 0 \\ 0 & -\frac{a_{32}+a_{33}}{a_{21}} & 1 \end{pmatrix}, \quad A_1 = \begin{pmatrix} -a_{11} & a_{12} + \frac{a_{13}(a_{32}+a_{33})}{a_{21}} & a_{13} \\ a_{21} & -a_{21} & 0 \\ 0 & m_1 & -a_{33} \end{pmatrix},$$

where $m_1 = \frac{a_{33}(a_{21}-a_{32}-a_{33})}{a_{21}}$. By the value of w_1 , the relevant discussion is divided into two subcases:

(i).
$$m_1 \neq 0$$
, (ii). $m_1 = 0$.

Case (i). If $m_1 \neq 0$, in view of the method introduced in Zhou et al. [28], it is assumed that $B_1 = H_1A_1H_1^{-1}$, where the standardized transformation matrix is

$$H_1 = \begin{pmatrix} a_{21}m_1 & -(a_{21}+a_{33})m_1 & a_{33}^2 \\ 0 & m_1 & -a_{33} \\ 0 & 0 & 1 \end{pmatrix}.$$
(4.8)

By direct calculation, one obtains

$$B_1 = \begin{pmatrix} -r_1 & -r_2 & -r_3 \\ 1 & 0 & 0 \\ 0 & 1 & 0 \end{pmatrix},$$

where r_1, r_2 and r_3 are the same as above. Furthermore, one can equivalently transform Eq. (4.7) into

$$(H_1J_1)M_1^2(H_1J_1)^{\tau} + B_1[(H_1J_1)\Sigma_1(H_1J_1)^{\tau}] + [(H_1J_1)\Sigma_1(H_1J_1)^{\tau}]B_1^{\tau} = 0$$

Letting $\Theta_0 = \varrho_1^{-2}(H_1J_1)\Sigma_1(H_1J_1)^{\tau}$, where $\varrho_1 = a_{21}m_1\sigma_1$, we obtain

 $G_0^2 + B_1 \Theta_0 + \Theta_0 B_1^\tau = 0.$

Noting that A has all negative real-part eigenvalues, then B_1 is a standard R_1 matrix. By Lemma 2.3, this means that Σ_0 is positive definite, which takes the form

$$\Theta_{0} = \begin{pmatrix} \frac{r_{2}}{2(r_{1}r_{2}-r_{3})} & 0 & -\frac{1}{2(r_{1}r_{2}-r_{3})} \\ 0 & \frac{1}{2(r_{1}r_{2}-r_{3})} & 0 \\ -\frac{1}{2(r_{1}r_{2}-r_{3})} & 0 & \frac{r_{1}}{2r_{3}(r_{1}r_{2}-r_{3})} \end{pmatrix}.$$
(4.9)

Therefore, $\Sigma_1 = \varrho_1^2 (H_1 J_1)^{-1} \Theta_0 [(H_1 J_1)^{-1}]^{\tau}$. Case (ii). If $m_1 = 0$, i.e., $a_{21} = a_{32} + a_{33}$, $\tilde{B}_1 = \tilde{H}_1 A_1 \tilde{H}_1^{-1}$ is defined, where another standardized transformation matrix \tilde{H}_1 and \tilde{B}_1 are obtained by

$$\widetilde{H}_{1} = \begin{pmatrix} a_{21} & -a_{21} & 0\\ 0 & 1 & 0\\ 0 & 0 & 1 \end{pmatrix}, \quad \widetilde{B}_{1} = \begin{pmatrix} -w_{1} & -w_{2} & -\xi_{1}\\ 0 & 1 & 0\\ 0 & 0 & -a_{33} \end{pmatrix},$$
(4.10)

where $w_1 = a_{11} + a_{21}$, $w_2 = a_{21}(a_{11} - a_{12} - a_{13})$, and ξ_1 is abbreviation. Obviously, \tilde{B}_1 is a standard R_2 matrix. Additionally, (4.7) can be equivalently transformed into

$$\begin{aligned} &(\widetilde{H}_1J_1)M_1^2(\widetilde{H}_1J_1)^{\tau} + \widetilde{B}_1[(\widetilde{H}_1J_1)\Sigma_1(\widetilde{H}_1J_1)^{\tau}] + [(\widetilde{H}_1J_1)\Sigma_1(\widetilde{H}_1J_1)^{\tau}]\widetilde{B}_1^{\tau} = 0. \\ \text{By letting } \Theta_1 &= (a_{21}\sigma_1)^{-2}(\widetilde{H}_1J_1)\Sigma_1(\widetilde{H}_1J_1)^{\tau}, \text{ it can be simplified as} \\ &G_0^2 + \widetilde{B}_1\Theta_1 + \Theta_1\widetilde{B}_1^{\tau} = 0. \\ \text{In view of Lemma } 2.4, \Theta_1 \text{ is described by} \end{aligned}$$

 $\Theta_1 = \begin{pmatrix} \frac{1}{2w_1} & 0 & 0\\ 0 & \frac{1}{2w_1w_2} & 0\\ 0 & 0 & 0 \end{pmatrix}.$

(4.11)

(4.12)

Hence, $\Sigma_1 = a_{21}^2 \sigma_1^2 (\tilde{H}_1 J_1)^{-1} \Theta_1 [(\tilde{H}_1 J_1)^{-1}]^{\tau}$. **Step 2.** Consider the algebraic equation $M_2^2 + A\Sigma_2 + \Sigma_2 A^{\tau} = 0.$

For the corresponding elimination matrix J_2 , J_3 , $A_2 = J_2 A J_2^{-1}$ is defined, where J_2 , J_3 and A_2 are obtained by

$$J_{2} = \begin{pmatrix} 0 & 1 & 0 \\ 0 & 0 & 1 \\ 1 & 0 & 0 \end{pmatrix}, \quad J_{3} = \begin{pmatrix} 1 & 0 & 0 \\ 0 & 1 & 0 \\ 0 & \frac{a_{12}}{a_{32}} & 1 \end{pmatrix}, \quad A_{2} = \begin{pmatrix} -a_{21} & -\frac{a_{12}a_{21}}{a_{32}} & a_{21} \\ -a_{32} & -a_{33} - a_{12} - \frac{a_{12}a_{33}}{a_{32}} & a_{32} + a_{33} \\ 0 & m_{2} & -a_{11} + a_{12} + \frac{a_{12}a_{33}}{a_{32}} \end{pmatrix},$$

where $m_2 = a_{13} + \frac{a_{12}(a_{11}-a_{33})}{a_{32}} - \frac{a_{12}^2(a_{32}+a_{33})}{a_{32}^2}$. Similarly, the following two sub-conditions are considered:

(1).
$$m_2 \neq 0$$
, (2). $m_2 = 0$

Case (1). If $m_2 \neq 0$, let $B_2 = H_2 A_2 H_2^{-1}$, where the relevant standardized transformation matrix

$$H_{2} = \begin{pmatrix} -a_{32}m_{2} & -(a_{11}+a_{33})m_{2} & (a_{11}-a_{12}-\frac{a_{12}a_{33}}{a_{32}})^{2} + m_{2}(a_{32}+a_{33}) \\ 0 & m_{2} & -a_{11}+a_{12}+\frac{a_{12}a_{33}}{a_{32}} \\ 0 & 0 & 1 \end{pmatrix}.$$
(4.13)

In fact, one still derives

$$B_2 = B_1 = \begin{pmatrix} -r_1 & -r_2 & -r_3 \\ 1 & 0 & 0 \\ 0 & 1 & 0 \end{pmatrix},$$

which means that B_2 is also a standard R_1 matrix. By letting $\Theta_2 = \varrho_2^{-2} (H_2 J_2) \Sigma_2 (H_2 J_2)^{\tau}$, where $\varrho_2 = -a_{32} m_2 \sigma_2$, (4.12) is then equivalent to the following equation:

 $G_0^2 + B_2 \Theta_2 + \Theta_2 B_2^{\tau} = 0.$

By Lemma 2.3 and the result of A having all negative real-part eigenvalues again, it can be shown that

$$\Theta_2 = \Theta_0 = \begin{pmatrix} \frac{r_2}{2(r_1r_2 - r_3)} & 0 & -\frac{1}{2(r_1r_2 - r_3)} \\ 0 & \frac{1}{2(r_1r_2 - r_3)} & 0 \\ -\frac{1}{2(r_1r_2 - r_3)} & 0 & \frac{r_1}{2r_3(r_1r_2 - r_3)} \end{pmatrix}.$$

In other words, $\Sigma_2 = \varrho_2^2 (H_2 J_3 J_2)^{-1} \Theta_0 [(H_2 J_3 J_2)^{-1}]^{\tau}$.

Case (2). If $m_2 = 0$, let $\tilde{B}_2 = \tilde{H}_2 A_2 \tilde{H}_2^{-1}$, where the corresponding standardized transformation matrix \tilde{H}_2 and \tilde{B}_2 are given by

$$\widetilde{H}_2 = \begin{pmatrix} -a_{32} & -a_{33} - a_{12} - \frac{a_{12}a_{33}}{a_{32}} & a_{32} + a_{33} \\ 0 & 1 & 0 \\ 0 & 0 & 1 \end{pmatrix}, \quad \widetilde{B}_2 = \begin{pmatrix} -w_3 & -w_4 & -\xi_2 \\ 0 & 1 & 0 \\ 0 & 0 & -a_{11} + a_{12} + \frac{a_{12}a_{33}}{a_{32}} \end{pmatrix},$$

where $w_3 = a_{12} + a_{21} + a_{33} + \frac{a_{12}a_{33}}{a_{32}}$, $w_4 = \frac{a_{21}a_{32}a_{33}(a_{11}-a_{12}-a_{13})}{a_{32}(a_{11}-a_{12})-a_{12}a_{33}}$, and ξ_2 is also shorthand. Similarly, \tilde{B}_2 is a standard R_2 matrix. By defining $\Theta_3 = (a_{32}\sigma_2)^{-2}(\tilde{H}_2J_2)\Sigma_2(\tilde{H}_2J_2)^{\tau}$, (4.12) is then equivalent to

$$G_0^2 + B_2 \Theta_3 + \Theta_3 B_2^\tau = 0.$$

According to Lemma 2.4, Θ_1 takes the form

$$\Theta_3 = \begin{pmatrix} \frac{1}{2w_3} & 0 & 0\\ 0 & \frac{1}{2w_3w_4} & 0\\ 0 & 0 & 0 \end{pmatrix}.$$
(4.14)

Then, $\Sigma_2 = a_{32}^2 \sigma_2^2 (\widetilde{H}_2 J_3 J_2)^{-1} \Theta_3 [(\widetilde{H}_2 J_3 J_2)^{-1}]^{\tau}$. **Step 3.** For the following algebraic equation,

 $M_3^2 + A\Sigma_3 + \Sigma_3 A^\tau = 0,$

and for the following elimination matrix J_3 , let $A_3 = J_4 A J_4^{-1}$, where J_4 and A_3 are given by

$$J_4 = \begin{pmatrix} 0 & 0 & 1 \\ 1 & 0 & 0 \\ 0 & 1 & 0 \end{pmatrix}, \quad A_3 = \begin{pmatrix} -a_{33} & a_{32} + a_{33} & -a_{32} \\ a_{13} & a_{12} & -a_{11} \\ 0 & a_{21} & -a_{21} \end{pmatrix}$$

Hence, (4.15) can be equivalently transformed into

$$J_4 M_3^2 J_4^{\tau} + B_3 \Theta_4 + \Theta_4 B_3^{\tau} = 0, \tag{4.16}$$

where $\Theta_4 = J_4 \Sigma_3 J_4^{\tau}$. Similarly, the proof is divided into two subcases by the value of a_{13} . Case (I). If $a_{13} \neq 0$, consider the corresponding standardized transformation matrix

$$H_3 = \begin{pmatrix} a_{13}a_{21} & -a_{21}(a_{11}+a_{21}) & a_{21}(a_{12}+a_{21}) \\ 0 & a_{21} & -a_{21} \\ 0 & 0 & 1 \end{pmatrix}.$$

(4.15)

Letting $B_3 = H_3A_3H_3^{-1}$, that $B_3 = B_1$ is still derived. Hence, an equivalent algebraic equation of (4.16) is described as follows:

$$(H_{3}J_{4})M_{3}^{2}(H_{3}J_{4})^{\tau} + B_{3}[(H_{3}J_{4})\Sigma_{3}(H_{3}J_{4})^{\tau}] + [(H_{3}J_{4})\Sigma_{3}(H_{3}J_{4})^{\tau}]B_{3}^{\tau} = 0.$$

Denoting $\Theta_{5} = \varrho_{3}^{-2}(H_{3}J_{4})\Sigma_{3}(H_{3}J_{4})^{\tau}$, where $\varrho_{3} = a_{13}a_{21}\sigma_{3}$, the last equation can be also simplified as
 $G_{0}^{2} + B_{1}\Theta_{5} + \Theta_{5}B_{1}^{\tau} = 0.$

Similarly, one obtains

$$\Theta_5 = \Theta_0 = \begin{pmatrix} \frac{r_2}{2(r_1r_2 - r_3)} & 0 & -\frac{1}{2(r_1r_2 - r_3)} \\ 0 & \frac{1}{2(r_1r_2 - r_3)} & 0 \\ -\frac{1}{2(r_1r_2 - r_3)} & 0 & \frac{r_1}{2r_3(r_1r_2 - r_3)} \end{pmatrix}.$$

Consequently, $\Sigma_3 = \varrho_3^2 (H_3 J_4)^{-1} \Theta_0 [(H_3 J_4)^{-1}]^{\tau}$. Case (II). If $a_{21} = 0$, then A_3 is a standard R_3 matrix. Noting that $J_4 M_3^2 J_4^{\tau} = \sigma_3^2 G_0^2 = \text{diag}(\sigma_3^2, 0, 0)$, by Lemma 2.5, one can obtain

$$\Theta_4 = J_4 \Sigma_3^2 J_4^{\tau} = \begin{pmatrix} \frac{\sigma_3^2}{2a_{33}} & 0 & 0\\ 0 & 0 & 0\\ 0 & 0 & 0 \end{pmatrix}.$$

Based on $a_{33} > 0$, then Θ_4 is a positive semi-definite matrix, which means that $\Sigma_3 = J_4^{-1} \Theta_4 (J_4^{-1})^{\tau}$. In summary, the special form of Σ is divided into eight cases by the different values of m_1 , m_2 and a_{13} , which is shown in Theorem 4.1. Finally, in view of the relation of systems (4.1) and (4.3), the stationary distribution $\overline{\omega}(\cdot)$ around E^* then has a unique log-normal probability density function

$$\Phi(S,V,I) = (2\pi)^{-\frac{3}{2}} |\Sigma|^{-\frac{1}{2}} e^{-\frac{1}{2}(\ln \frac{S}{S^*}, \ln \frac{V}{V^*}, \ln \frac{I}{I^*})\Sigma^{-1}(\ln \frac{S}{S^*}, \ln \frac{V}{V^*}, \ln \frac{I}{I^*})^{\tau}}.$$

Therefore, this completes the proof. \Box

Remark 3. If $\mathscr{P}_0^{\epsilon} > 1$, Theorem 4.1 shows that the stationary distribution $\varpi(\cdot)$ around E^* has the unique log-normal density function $\Phi(S, V, I)$. This reflects the stochastic permanence of system (1.3) from one side. In addition, that $\mathscr{R}_0^s = \mathscr{R}_0^c = \mathscr{R}_0$ if $\sigma_i = 0$ (i = 1, 2, 3) is obtained.

5. Extinction of system (1.3)

As is known, all of the properties of disease persistence of system (1.3) are reflected by Theorems 3.1 and 4.1. For a comprehensive study, a simple extinction result of system (1.3) is described by the following Theorem 5.1.

Theorem 5.1. For any initial value $(S(0), V(0), I(0)) \in \mathbb{R}^3_+$, if $\mathscr{R}^d_0 = \frac{\beta}{\mu + \alpha + \delta + \frac{\sigma_1^2}{2}} < 1$, then the solution (S(t), V(t), I(t)) of system (1.3) follows:

$$\limsup_{t \to +\infty} \frac{\ln I(t)}{t} \le \left(\mu + \alpha + \delta + \frac{\sigma_3^2}{2}\right) \left(\mathscr{R}_0^d - 1\right) < 0, \quad a.s.,$$
(5.1)

which means that the epidemic of system (1.3) will go to extinction with probability 1 (a.s.).

Proof. Employing Itô's formula to $\ln I(t)$, one obtains

$$d\ln I(t) = \left[\frac{\beta S(t)}{N(t)} - \left(\mu + \alpha + \delta + \frac{\sigma_3^2}{2}\right)\right] dt + \sigma_3 dB_3(t).$$
(5.2)

Integrating from 0 to t and dividing by t on both sides of (5.1), it can be seen that

$$\frac{\ln I(t)}{t} \leq \frac{\ln I(0)}{t} + \frac{1}{t} \int_{0}^{t} \left[\frac{\beta S(u)}{N(u)} - \left(\mu + \alpha + \delta + \frac{\sigma_{3}^{2}}{2}\right) \right] du + \frac{\int_{0}^{t} \sigma_{3} dB_{3}(u)}{t} \\
\leq \frac{\ln I(0)}{t} + \frac{1}{t} \int_{0}^{t} \left[\beta - \left(\mu + \alpha + \delta + \frac{\sigma_{3}^{2}}{2}\right) \right] du + \frac{\int_{0}^{t} \sigma_{3} dB_{3}(u)}{t} \\
= \frac{\ln I(0)}{t} + \left(\mu + \alpha + \delta + \frac{\sigma_{3}^{2}}{2}\right) \left(\mathscr{R}_{0}^{d} - 1 \right) + \frac{\int_{0}^{t} \sigma_{3} dB_{3}(u)}{t}.$$
(5.3)

Next, by the strong law of large numbers [1], one derives

$$\lim_{t \to +\infty} \frac{\int_0^t \sigma_3 dB_3(u)}{t} = 0, \quad a.s.$$
(5.4)

Taking the superior limit of $t \to +\infty$ on both sides of (5.3), the assertion (5.1) can then be obtained by (5.4). Moreover, from the expressions of \mathscr{R}_0^s and \mathscr{R}_0^s , one can obtain that $\mathscr{R}_0^d \leq \mathscr{R}_0^s$.

Consequently, the proof of Theorem 5.1 is confirmed. \Box

Parameters	Description	Unit	Value	Source
Λ	Recruitment rate of population	per day	≥ 0.5	[38,39]
β	Transmission rate of susceptible individuals	per day	[0.390,0.432]	[13]
μ	Natural death rate of population	per day	$[2.74, 6.85] \times 10^{-5}$	[47],CSZ data
α	Disease mortality of infected people	per day	$\frac{1}{0.6\times 365}$	[13]
δ	Recovery rate	None	[0.01,0.2]	Estimated
γ	Immune loss rate of vaccinated individuals	None	0.2	[37]
θ	Vaccination rate of susceptible individuals	None	[0.371,0.436]	[13]

List of biological parameters of system (1.3)

6. Simulations and parameter analyses

Table 1

In this section, by means of the well-known higher-order method developed by Milstein [46], the corresponding discretization equation of system (1.3) is obtained in the form

$$\begin{cases} S^{k+1} = S^k + \left[\Lambda - (\mu + \vartheta)S^k - \frac{\beta S^k I^k}{S^k + V^k + I^k} + \gamma V^k + \delta I^k\right] \Delta t + \frac{\sigma_1^2}{2} S^k (\xi_k^2 - 1) \Delta t + \sigma_1 S^k \sqrt{\Delta t} \xi_k \\ V^{k+1} = V^k + \left[\vartheta S^k - (\mu + \gamma)V^k\right] \Delta t + \frac{\sigma_2^2}{2} V^k (\eta_k^2 - 1) \Delta t + \sigma_2 V^k \sqrt{\Delta t} \eta_k \\ I^{k+1} = I^k + \left[\frac{\beta S^k I^k}{S^k + V^k + I^k} - (\mu + \alpha + \delta)I^k\right] \Delta t + \frac{\sigma_3^2}{2} I^k (\zeta_k^2 - 1) \Delta t + \sigma_3 I^k \sqrt{\Delta t} \zeta_k, \end{cases}$$

$$\tag{6.1}$$

where the time increment $\Delta t > 0$, and ξ_k , η_k , $and\zeta_k$ are three independent Gaussian random variables that follow the distribution N(0, 1) for k = 1, 2, ..., n. Furthermore, (S^k, V^k, I^k) is the corresponding value of the *k*th iteration of the discretization equation. From AI-Darabsah [13], Zhao and Jiang [37], Liu et al. [38], Zhang and Jiang [39], Arino et al. [47], and the detailed data of the Central Statistical Office of Zimbabwe (CSZ), the corresponding realistic statistics of system (1.3) are shown in Table 1. Next, several empirical examples are provided to focus on the following five aspects.

- (i) The existence of the ergodic stationary distribution of system (1.3) while $\Re_0^s > 1$.
- (ii) The exact expression and verification of the unique log-normal density function for the stationary distribution under $\mathscr{R}_0^s > 1$.
- (iii) The influence of random fluctuations on the disease persistence of system (1.3).
- (iv) The effects of the main parameters of system (1.3) on the disease dynamics.
- (v) The corresponding dynamical behavior of system (1.3) if $\mathscr{R}_0^d < 1$.

6.1. Dynamical behavior of system (1.3) if $\mathscr{R}_0^s > 1$

Example 6.1. By Table 1, letting the environmental noise intensities $(\sigma_1, \sigma_2, \sigma_3) = (0.0008, 0.0004, 0.0008)$ and main parameters $(\Lambda, \beta, \mu, \alpha, \delta, \gamma, \vartheta) = (0.8, 0.4, 3 \times 10^{-5}, 0.00457, 0.05, 0.2, 0.4)$, one then obtains

æ2.

$$\mathscr{R}_{0} = \frac{\beta(\mu+\gamma)}{(\mu+\gamma+\vartheta)(\mu+\alpha+\delta)} = 2.44 > 1, \quad \mathscr{R}_{0}^{s} = \frac{\mu\beta\left(\mu+\gamma+\frac{\sigma_{2}^{2}}{2}\right)}{\left[\left(\mu+\frac{\sigma_{1}^{2}}{2}\right)\left(\mu+\gamma+\frac{\sigma_{2}^{2}}{2}\right)+\vartheta\left(\mu+\frac{\sigma_{2}^{2}}{2}\right)\right]\left(\mu+\alpha+\delta+\frac{\sigma_{3}^{2}}{2}\right)} = 2.43 > 1.$$

It follows from Theorem 3.1 that system (1.3) admits a unique ergodic stationary distribution $\varpi(\cdot)$. The left-hand column of Fig. 1 can be seen to validate it. By Theorem 4.1, the stationary distribution $\varpi(\cdot)$ around the quasi-endemic equilibrium E^* has a unique log-normal density function $\Phi(S, V, I)$. Moreover, it is calculated that

 $m_1=0.0246\neq 0, \quad m_2=-27.8515\neq 0, \quad a_{13}=0.1196\neq 0,$

which means

$$\begin{split} \Sigma &= \varrho_1^2 (H_1 J_1)^{-1} \Theta_0 [(H_1 J_1)^{-1}]^{\tau} + \varrho_2^2 (H_2 J_3 J_2)^{-1} \Theta_0 [(H_2 J_3 J_2)^{-1}]^{\tau} + \varrho_3^2 (H_3 J_4)^{-1} \Theta_0 [(M_3 J_3)^{-1}]^{\tau} \\ &= 10^{-4} \times \begin{pmatrix} 0.3524 & 0.3498 & 0.3947 \\ 0.3498 & 0.3538 & 0.3882 \\ 0.3947 & 0.3882 & 0.4970 \end{pmatrix}. \end{split}$$

By direct calculation, one can obtain that $E^* = (S^*, V^*, I^*) = (40.0428, 173.2562, 80.0736)$. Then, the corresponding marginal density functions of S(t), V(t) and I(t) are separately given as follows.

(1).
$$P_1(S) = \frac{\partial \Phi}{\partial S} = 67.204e^{-14188.4(\ln S - 3.69)}$$
, (ii). $P_2(V) = \frac{\partial \Phi}{\partial V} = 67.07e^{-14132.2(\ln V - 5.15)}$,

(iii).
$$P_3(I) = \frac{\partial \Phi}{\partial I} = 56.59e^{-10060.4(\ln I - 4.38)}$$
.

The curves of (i)-(iii) are shown in the right-hand column of Fig. 1. Obviously, this greatly illustrates Theorem 4.1 from the side.

Combining Remarks 3.1–4.1 and Theorem 5.1, one can derive that all random perturbations σ_1 , σ_2 , and σ_3 have a critical influence on the dynamical behavior of system (1.3). Therefore, the corresponding parameter analyses of the above three white noises are shown by Example 6.2.



Fig. 1. Left-hand column shows simulation of compartments S(t), V(t), and I(t) in deterministic system (1.1) and stochastic system (1.3) with noise intensities (σ_1 , σ_2 , σ_3) = (0.0008, 0.0004, 0.0008) and main parameters (Λ , β , μ , α , δ , γ , ϑ) = (0.8, 0.4, 3 × 10⁻⁵, 0.00457, 0.05, 0.2, 0.4), respectively. Right-hand column shows frequency histogram and corresponding marginal density function curves of individuals *S*, *V*, and *I*.



Fig. 2. Corresponding simulation of partial compartments S(t) and I(t) of stochastic system (1.3) under noise intensities $(\sigma_1, \sigma_2, \sigma_3) = (0.0008, 0.0004, 0.0008), (0.008, 0.0004, 0.0008), (0.0008, 0.004, 0.0008)$ and (0.0008, 0.0004, 0.008), respectively. Other fixed parameters: $(\Lambda, \beta, \mu, \alpha, \delta, \gamma, \vartheta) = (0.8, 0.4, 3 \times 10^{-5}, 0.00457, 0.05, 0.2, 0.4).$

6.2. Impact of random noises σ_i (i = 1, 2, 3) on disease extinction and the existence of stationary distribution

Example 6.2. One chooses the epidemiological parameters $(\Lambda, \beta, \mu, \alpha, \delta, \gamma, \vartheta) = (0.8, 0.4, 3 \times 10^{-5}, 0.00457, 0.05, 0.2, 0.4)$ and considers the following four subcases of stochastic perturbations:

(i). $(\sigma_1, \sigma_2, \sigma_3) = (0.0008, 0.0004, 0.0008),$ (ii). $(\sigma_1, \sigma_2, \sigma_3) = (0.008, 0.0004, 0.0008),$

(iii). $(\sigma_1, \sigma_2, \sigma_3) = (0.0008, 0.004, 0.0008),$ (iv). $(\sigma_1, \sigma_2, \sigma_3) = (0.0008, 0.0004, 0.008).$

First, it should be pointed out that the above four subcases (i)–(iv) all guarantee the existence of a stationary distribution, which has an ergodicity property. For convenience and simplicity, only the population intensities of susceptible and infected individuals are focused on, which are presented in subfigures (2-1) and (2-2) of Fig. 2, respectively. By only increasing the perturbation intensities of the vaccinated individuals (or infected individuals), i.e., the larger σ_2 (or σ_3), then the disease infection will be effectively inhibited. In contrast, by only increasing the perturbation intensity of the susceptible individuals, a great destabilizing influence on the population numbers of *S* and *I* manifests.



Fig. 3. Corresponding population numbers of solution (S(t), V(t), I(t)) to system (1.3) with transmission rates of $\beta = 0.39$, 0.40, 0.41, and 0.42, respectively. Other given parameters: (Λ , μ , α , δ , γ , ϑ) = (0.8, 3 × 10⁻⁵, 0.00457, 0.128, 0.2, 0.4) and (σ_1 , σ_2 , σ_3) = (0.0008, 0.0004, 0.0008).



Fig. 4. Corresponding simulation of solution (*S*(*t*), *V*(*t*), *I*(*t*)) to system (1.3) with vaccination rate $\vartheta = 0.371$, 0.386, 0.401, and 0.416, respectively. Other fixed parameters: $(\Lambda, \beta, \mu, \alpha, \delta, \gamma) = (0.8, 0.4, 3 \times 10^{-5}, 0.00457, 0.128, 0.2)$ and $(\sigma_1, \sigma_2, \sigma_3) = (0.0008, 0.0004, 0.0008)$.

Next, by Zhu et al. [25], Jia et al. [26], the impact of the main parameters of system (1.3) on the individual decision-making behavior is studied. From the expressions of \mathscr{R}_0^s and \mathscr{R}_0^d , the disease persistence and extinction of system (1.3) are critically affected by the transmission rate β and vaccination rate ϑ . Thus, the following Examples 6.3 and 6.4 will reveal the impact. In addition, the corresponding effect of the recruitment rate Λ on the dynamical behavior of system (1.3) is also shown in Example 6.5.

6.3. Impact of transmission rate β on dynamics of system (1.3)

Example 6.3. Choosing the epidemiological parameters $(\Lambda, \mu, \alpha, \delta, \gamma, \vartheta) = (0.8, 3 \times 10^{-5}, 0.00457, 0.128, 0.2, 0.4)$ and random noises $(\sigma_1, \sigma_2, \sigma_3) = (0.0008, 0.0004, 0.0008)$ and considering the subcases of transmission rate $\beta = 0.39$, 0.40, 0.41 and 0.42, the corresponding numbers of the solution (S(t), V(t), I(t)) to system (1.3) are described in Fig. 3. Clearly, a small transmission rate can lead to reduction of disease infection and even elimination, such as $\beta \leq 0.39$ per day.

6.4. Impact of vaccination rate ϑ on dynamics of system (1.3)

Example 6.4. Assuming that the parameters $(\Lambda, \beta, \mu, \alpha, \delta, \gamma) = (0.8, 0.4, 3 \times 10^{-5}, 0.00457, 0.128, 0.2)$ and stochastic perturbations $(\sigma_1, \sigma_2, \sigma_3) = (0.0008, 0.0004, 0.0008)$, for the corresponding subcases of vaccination rate $\vartheta = 0.371$, 0.386, 0.401, and 0.416, the corresponding solutions (S(t), V(t), andI(t)) to system (1.3) are shown in Fig. 4. Similarly, a small vaccination rate can control the disease infection more effectively than a large one.



Fig. 5. Corresponding population intensities of individuals *S*, *V*, and *I* of system (1.3) with recruitment rate $\Lambda = 0.7$, 0.8, 0.9, and 1.0, respectively. Other given parameters: $(\beta, \mu, \alpha, \delta, \gamma, \vartheta) = (0.4, 3 \times 10^{-5}, 0.00457, 0.128, 0.2, 0.4)$ and $(\sigma_1, \sigma_2, \sigma_3) = (0.0008, 0.0004, 0.0008)$.



Fig. 6. Corresponding population numbers of solution (S(t), V(t), I(t)) to system (1.3) with random perturbations (σ_1 , σ_2 , σ_3) = (0.01, 0.01, 0.78) and main parameters (Λ , β , μ , α , δ , γ , ϑ) = (0.8, 0.4, 3 × 10^{-5}, 0.00457, 0.13, 0.2, 0.38).

6.5. Impact of recruitment rate Λ on dynamics of system (1.3)

Example 6.5. Letting the dynamical parameters be $(\beta, \mu, \alpha, \delta, \gamma, \vartheta) = (0.4, 3 \times 10^{-5}, 0.00457, 0.128, 0.2, 0.4)$ and stochastic fluctuations be $(\sigma_1, \sigma_2, \sigma_3) = (0.0008, 0.0004, 0.0008)$, and considering the sub-conditions of recruitment rate $\Lambda = 0.7$, 0.408, 0.9, and 1.0, the corresponding intensities of the compartments *S*, *V*, and *I* of system (1.3) are reflected in Fig. 5. Obviously, as the parameter Λ increases to 1 from 0.7, the spread and infection of an epidemic can be effectively controlled by the small recruitment rate.

6.6. Dynamical behaviors of system (1.3) under $\mathscr{R}_0^d < 1$

Example 6.6. Considering the stochastic noises $(\sigma_1, \sigma_2, \sigma_3) = (0.01, 0.01, 0.78)$ and main parameters $(\Lambda, \beta, \mu, \alpha, \delta, \gamma, \vartheta) = (0.8, 0.4, 3 \times 10^{-5}, 0.00457, 0.13, 0.2, 0.38)$, one can then obtain

$$\mathscr{R}_{0} = \frac{\beta(\mu + \gamma)}{(\mu + \gamma + \vartheta)(\mu + \alpha + \delta)} = 1.0249 > 1, \quad \mathscr{R}_{0}^{d} = \frac{\beta}{\mu + \alpha + \delta + \frac{\sigma_{3}^{2}}{2}} = 0.9116 < 1, \quad \mathscr{R}_{0}^{s} = 0.1179 < 1.$$

By Theorem 3.1, one cannot derive the existence of the ergodic stationary distribution of system (1.3). In contrast, it follows from Theorem 5.1 that the disease of stochastic system (1.3) will be extinct in a long term. In addition, the deterministic model (1.1) has a globally asymptotically stable endemic equilibrium E^+ . On the one hand, these results validate the fact that large white noises lead to disease elimination from the side. On the other hand, the large random fluctuation σ_3 (*i.e.*, $\frac{\sigma_3}{\sigma_1} = \frac{\sigma_3}{\sigma_2} = 78 >> 1$) indicates that it is necessary to isolate and control the infected individuals during the outbreak of an epidemic. These results are verified by Fig. 6.

For epidemiological study, combining the above numerical simulations and parameter analyses, several reasonable and effective measures to reduce the threat of infectious diseases to human life, and even eliminate the epidemic, are provided. The special approaches are the following.

(i) Several reasonable policies of joint prevention and control are implemented to reduce the population mobility in differential risk epidemic areas. Then, the small recruitment rate Λ may lead to the elimination of disease (see Fig. 5).

(ii) Controlling the activities of the susceptible individuals in highly pathogenic areas to decrease the contact rate of population. Hence, $\beta \to 0^+$ can be guaranteed, which means $\mathscr{R}_0^s < 1$ and $\mathscr{R}_0^d < 1$ (see Fig. 3).

(iii) Developing several effective vaccines and carrying out other prophylactic measures to improve the immune rate of disease (see Fig. 4).

7. Conclusions and result discussions

7.1. Conclusions

The corresponding theoretical results of this paper are the following. (i) By Theorem 3.1, system (1.3) admits a unique ergodic stationary distribution $\overline{\omega}(\cdot)$ under

$$\mathscr{R}_{0}^{s} = \frac{\mu\beta\left(\mu + \gamma + \frac{\sigma_{2}^{2}}{2}\right)}{\left[\left(\mu + \frac{\sigma_{1}^{2}}{2}\right)\left(\mu + \gamma + \frac{\sigma_{2}^{2}}{2}\right) + \vartheta\left(\mu + \frac{\sigma_{2}^{2}}{2}\right)\right]\left(\mu + \alpha + \delta + \frac{\sigma_{3}^{2}}{2}\right)}$$

(ii) By taking the effect of random perturbations into account, a quasi-endemic equilibrium E^* related to E^+ is defined while $\mathscr{R}_0^c =$

 $\frac{\beta(\mu+\gamma+\frac{\sigma_2^2}{2})}{(\mu+\gamma+\epsilon+\frac{\sigma_2^2}{2})(\mu+\alpha+\delta+\frac{\sigma_1^2}{2})} > 1. \text{ In view of the expressions of } \mathscr{R}_0^c \text{ and } \mathscr{R}_0^s, \text{ it is further proved that the stationary distribution } \varpi(\cdot) \text{ around } E^* \text{ has a log-normal density function in the following form:}$

 $\Phi(S, V, I) = (2\pi)^{-\frac{3}{2}} |\Sigma|^{-\frac{1}{2}} e^{-\frac{1}{2}(\ln \frac{S}{S^{*}}, \ln \frac{V}{V^{*}}, \ln \frac{I}{I^{*}})\Sigma^{-1}(\ln \frac{S}{S^{*}}, \ln \frac{V}{V^{*}}, \ln \frac{I}{I^{*}})^{*}}.$

where the special form of Σ is shown in Theorem 4.1. (iii) The disease of system (1.3) will go to extinction with probability 1 if \mathscr{R}_0^d = $\frac{\beta}{\mu + \alpha + \delta + \frac{\sigma_3^2}{2}}$ < 1. The above results (i) and (ii) reflect the stochastic persistence and ergodicity of the epidemic. Moreover, the corresponding

disease extinction of system (1.3) is described by result (iii).

7.2. Result discussions

In this paper, combining the great effect of vaccination and the unpredictability of environmental fluctuations in the real world, a stochastic SVIS infectious disease model with vaccination and standard incidence is the object of concern. Adopting the descriptions in [28-40], linear perturbation, which is the most intuitive assumption of a random effect, is similarly taken into consideration in this paper. Subsequently, several dynamical properties of stochastic system (1.3) are analyzed, such as the existence and uniqueness of a global positive solution, existence and ergodicity of a stationary distribution, and disease elimination. By comparison with the existing results ([28–40]), several highlights developed in the present study are detailed in the following two points.

• As is known, the endemic equilibrium and basic reproduction number, two important results of a deterministic epidemic, reflect disease permanence and elimination. Similarly, for the corresponding stochastic model, the existence of stationary distribution indicates the stochastic positive equilibrium state. In this paper, it is first proved that stochastic system (1.3) admits a unique ergodic stationary distribution under the critical value $\mathscr{R}_0^s > 1$. It should be pointed out that $\mathscr{R}_0^s > 1$ is a unified threshold for the disease persistence of systems (1.1) and (1.3). Moreover, the sufficient condition $\mathscr{R}_0^d < 1$ is obtained for the disease extinction of system (1.3). Both $\mathscr{R}_0^s > 1$ and $\mathscr{R}_0^d < 1$ reveal that the dynamical behavior of system (1.3) is critically affected by the random fluctuations, i.e., σ_1, σ_2 , and σ_3 . In view of the method of controlling variables and numerical simulations, this means that a large white noise leads to the disease eradication, while a small one guarantees stochastic permanence. In addition, by the main parameter analyses, several effective measures to stop the spread of an epidemic are provided.

• It is generally accepted that the existence of an ergodic stationary distribution incurs difficulty in studying more exact statistical properties. Hence, this paper is devoted to obtaining the corresponding probability density function for further dynamical investigation. The results of Zhou et al. [28] are further perfected and general solving theories of algebraic equations with respect to the three-dimensional Fokker-Planck equation are developed, which are described in Lemmas 2.5 and 2.6. By taking the effect of stochasticity into account again, the quasi-endemic equilibrium E^* corresponding to the endemic equilibrium E^+ is defined. For practical application, the exact expression of the log-normal three-dimensional density function $\Phi(S, V, I)$ of system (1.3) is given. Furthermore, it is worth mentioning that the methods and theories developed herein are still suitable for the case of the diffusion matrix M being positive semi-definite, such as delay stochastic differential equations [32,48].

Several remaining issues are now proposed and analyzed. First, by virtue of the limitation of the present mathematical approaches for epidemiological dynamics, a value gap exists between \mathscr{R}_0^s and \mathscr{R}_0^d , and it is unfortunate that difficulty is encountered in obtaining the most precise threshold for disease extinction and persistence. Second, the impact of telegraph noises and periodicity on the dynamics of system (1.3) should also be studied; the reader is referred to [30,34,45,49,50]. These problems are expected to be studied and solved as planned future work.

Declaration of Competing Interest

We declare that we do not have any commercial or associative interest that represents a conflict of interest in connection with the work submitted.

CRediT authorship contribution statement

Baoquan Zhou: Validation, Software, Formal analysis, Writing - original draft, Writing - review & editing. Daging Jiang: Conceptualization, Investigation, Methodology, Writing - review & editing. Yucong Dai: Methodology, Formal analysis, Writing - original draft, Writing - review & editing. **Tasawar Hayat:** Conceptualization, Writing - review & editing. **Ahmed Alsaedi:** Investigation, Writing - original draft, Writing - review & editing.

Acknowledgements

This work was supported by the National Natural Science Foundation of China (Grant No. 11871473) and Shandong Provincial Natural Science Foundation (Grant Nos. ZR2019MA010 and ZR2019MA006).

Appendix A.

(I) (Proof of Lemma 2.5): Consider the algebraic equation $G_1^2 + C_0 \Upsilon_3 + \Upsilon_3 C_0^{\tau} = 0$, where Υ_3 is a symmetric matrix. Letting $\Upsilon_3 := (\kappa_{ij})_{3\times 3}$, by direct calculation one has

$$\Upsilon_3 = \begin{pmatrix} \kappa_{11} & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{pmatrix},\tag{A.1}$$

where $\kappa_{11} = -\frac{a_0^2}{2c_{11}}$. If $c_{11} < 0$, then it means that Υ_3 is a positive semi-definite matrix. The proof is then completed. **(II)** (Proof of Lemma 2.6): Denote

$$\Pi_1 = \begin{pmatrix} 1 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{pmatrix}, \ \Pi_2 = \begin{pmatrix} 0 & 0 & 0 \\ 0 & 1 & 0 \\ 0 & 0 & 0 \end{pmatrix}, \ \Pi_3 = \begin{pmatrix} 0 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 1 \end{pmatrix}.$$

Let Σ_i (*i* = 1, 2, 3) be the solutions of the following algebraic equations, respectively:

 $\Pi_i + A\Sigma_i + \Sigma_i A^{\tau} = 0.$

Obviously, one can obtain

 $\Sigma_0 = \alpha_1^2 \Sigma_1 + \alpha_2^2 \Sigma_2 + \alpha_3^2 \Sigma_3.$

Before proving the positive definiteness of Σ_0 , the following two theories of matrix algebra should be described first.

 (T_1) . The positive definiteness of the matrix is not affected by the inverse congruence transformation.

 (T_2) . The similarity transformation does not change the eigenvalues of the matrix.

For convenience and simplicity, an important notation is introduced as follows. For the same dimensional square matrix A and B, define

 $A \succeq B$: A - B is at least a positive semi-definite matrix.

Given the above, it is easily derived that *A* is also positive definite if *B* is a positive definite matrix.

First, consider the following algebraic equation,

$$\Pi_1 + A\Sigma_1 + \Sigma_1 A^{\tau} = 0,$$

after which the relevant proof can be divided into the following two conditions:

$$(\mathscr{B}_1). \ a_{21} = a_{31} = 0, \ (\mathscr{B}_2). \ a_{21} \neq 0 \text{ or } a_{31} \neq 0.$$

Next, one must demonstrate that the elements a_{21} and a_{31} have the equivalent status in *A*. Let $\tilde{A} = F_1 A F_1^{-1} := (\tilde{a}_{ij})_{3 \times 3}$, where \tilde{A} and the invertible matrix F_1 are given by

$$F_1 = \begin{pmatrix} 1 & 0 & 0 \\ 0 & 0 & 1 \\ 0 & 1 & 0 \end{pmatrix}, \quad \widetilde{A} = \begin{pmatrix} a_{11} & a_{13} & a_{12} \\ a_{31} & a_{33} & a_{32} \\ a_{21} & a_{23} & a_{22} \end{pmatrix}$$

respectively. Hence, (A.2) can be equivalently transformed as

$$F_1 \Pi_1 F_1^{\tau} + \hat{A} (F_1 \Sigma_1 F_1^{\tau}) + (F_1 \Sigma_1 F_1^{\tau}) \hat{A}^{\tau} = 0.$$

By defining $\widetilde{\Pi}_1 = F_1 \Pi_1 F_1^{\tau}$, $\widetilde{\Sigma}_1 = F_1 \Sigma_1 F_1^{\tau}$, it can be noticed that

(i). $\widetilde{\Pi}_1 = \Pi_1$, (ii). $\widetilde{\Sigma}_1$ and Σ_1 have the same positive definiteness.

In addition, $\tilde{a}_{21} = a_{31}$, $\tilde{a}_{31} = a_{21}$. Therefore, the validation is completed. Namely, one must only discuss the following two cases, which are equivalent to conditions (\mathscr{B}_1) and (\mathscr{B}_2), respectively:

$$(\mathscr{C}_1). \ a_{21} = a_{31} = 0, \ (\mathscr{C}_2). \ a_{21} \neq 0.$$

Case (\mathscr{C}_1). If $a_{21} = a_{31} = 0$, by directly solving Eq. (A.2), one obtains

$$\Sigma_1 = \begin{pmatrix} -\frac{1}{2a_{11}} & 0 & 0\\ 0 & 0 & 0\\ 0 & 0 & 0 \end{pmatrix} := \Delta_{11}.$$

Since A has all negative real part eigenvalues, by the similarity invariance of $\psi_A(\lambda)$, it indicates that

 $\lambda^3 + r_1\lambda^2 + r_2\lambda + r_3 = \psi_A(\lambda) = (\lambda - a_{11})[\lambda^2 - (a_{22} + a_{33})\lambda + (a_{22}a_{33} - a_{23}a_{32})].$

(A.3)

(A.2)

Consequently, $\varphi_A(\lambda)$ has an eigenvalue $\lambda_1 = a_{11}$, which has a negative real part. By $a_{11} \in \mathbb{R}$, then $a_{11} < 0$. In other words, Δ_{11} is positive semi-definite. Moreover,

$$\Sigma_1 \succeq \Delta_{11}. \tag{A.4}$$

Case (\mathscr{C}_2). If $a_{21} \neq 0$, let $\omega_0 = a_{32} + \frac{a_{31}(a_{33} - a_{22})}{a_{21}} - \frac{a_{23}a_{31}^2}{a_{21}^2}$. • If $\omega_0 = 0$, which means $a_{21}(a_{21}a_{32} - a_{22}a_{31}) - a_{31}(a_{31}a_{23} - a_{21}a_{33}) = 0$, let $\widehat{A} = (F_3F_2)A(F_3F_2)^{-1}$, where $\begin{pmatrix} 1 & 0 & 0 \end{pmatrix}$ $\begin{pmatrix} a_{21} & a_{22} + \frac{a_{23}a_{31}}{a_{21}} & a_{23} \end{pmatrix}$

$$F_2 = \begin{pmatrix} 1 & 0 & 0 \\ 0 & 1 & 0 \\ 0 & -\frac{a_{31}}{a_{21}} & 1 \end{pmatrix}, \quad F_3 = \begin{pmatrix} a_{21} & a_{22} & a_{23} \\ 0 & 1 & 0 \\ 0 & 0 & 1 \end{pmatrix}.$$

Thus, (A.2) can be equivalently rewritten as

$$(F_{3}F_{2})\Lambda_{1}(F_{3}F_{2})^{\tau} + \hat{A}[(F_{3}F_{2})\Sigma_{1}(F_{3}F_{2})^{\tau}] + [(F_{3}F_{2})\Sigma_{1}(F_{3}F_{2})^{\tau}]\hat{A}^{\tau} = 0.$$
(A.5)

Denoting $\widehat{\Pi}_1 = (F_3F_2)\Pi_1(F_3F_2)^{\tau}$ and $\widehat{\Sigma}_1 = (F_3F_2)\Sigma_1(F_3F_2)^{\tau}$, one computes

$$\widehat{\Pi}_1 = \begin{pmatrix} a_{21}^2 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{pmatrix}, \ \widehat{A} = \begin{pmatrix} -\xi_1 & -\xi_2 & -\xi_3 \\ 1 & 0 & 0 \\ 0 & 0 & a_{33} - \frac{a_{23}a_{31}}{a_{21}} \end{pmatrix}, \ \widehat{\Sigma}_1 = \begin{pmatrix} \frac{a_{21}^2}{2\xi_1} & 0 & 0 \\ 0 & \frac{a_{21}^2}{2\xi_1\xi_2} & 0 \\ 0 & 0 & 0 \end{pmatrix},$$

where the parameters ξ_k (k = 1, 2, 3) can be obtained by (A.5). Because their sign is the only object of concern, they are omitted here. Furthermore, the characteristic polynomial $\psi_A(\lambda)$ follows from \widehat{A} that

$$\psi_A(\lambda) = \left(\lambda - a_{33} + \frac{a_{23}a_{31}}{a_{21}}\right)(\lambda^2 + \xi_1\lambda + \xi_2)$$

By the condition that A has all negative real part eigenvalues, it thus means that the equation $\lambda^2 + \xi_1 \lambda + \xi_2 = 0$ has two negative real part roots. By the Routh-Hurwitz stability criterion [43], it can be shown that

$$\xi_1 > 0, \quad \xi_2 > 0.$$
 (A.6)

In view of

$$\widehat{\Sigma}_{1} = \begin{pmatrix} \frac{a_{21}^{2}}{2\xi_{1}} & 0 & 0\\ 0 & \frac{a_{21}^{2}}{2\xi_{1}\xi_{2}} & 0\\ 0 & 0 & 0 \end{pmatrix} = \begin{pmatrix} \frac{a_{21}^{2}}{2\xi_{1}} & 0 & 0\\ 0 & 0 & 0\\ 0 & 0 & 0 \end{pmatrix} + \begin{pmatrix} 0 & 0 & 0\\ 0 & \frac{a_{21}^{2}}{2\xi_{1}\xi_{2}} & 0\\ 0 & 0 & 0 \end{pmatrix} := L_{1} + L_{2},$$

one hence obtains

$$\begin{split} \Sigma_{1} &= (F_{3}F_{2})^{-1}\widehat{\Sigma}_{1}[(F_{3}F_{2})^{-1}]^{\tau} = (F_{3}F_{2})^{-1}(L_{1}+L_{2})[(F_{3}F_{2})^{-1}]^{\tau} \\ &= (F_{3}F_{2})^{-1}L_{1}[(F_{3}F_{2})^{-1}]^{\tau} + (F_{3}F_{2})^{-1}L_{2}[(F_{3}F_{2})^{-1}]^{\tau} \\ &= \begin{pmatrix} \frac{1}{2\xi_{1}} & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{pmatrix} + (F_{3}F_{2})^{-1}L_{2}[(F_{3}F_{2})^{-1}]^{\tau} \\ &:= \Delta_{12} + (F_{3}F_{2})^{-1}L_{2}[(F_{3}F_{2})^{-1}]^{\tau}. \end{split}$$
(A.7)

By means of $\xi_1 > 0$ and $\xi_2 > 0$, it is derived that Δ_{12} , L_2 and $(F_3F_2)^{-1}L_2[(F_3F_2)^{-1}]^{\tau}$ are all positive semi-definite. It is then implied that $\Sigma_1 > \Delta_{12}$. (A.8)

• If $\omega_0 \neq 0$, let $\bar{A} = (F_4F_2)A(F_4F_2)^{-1}$, where the invertible matrix F_4 is given by

$$F_4 = \begin{pmatrix} a_{21}\omega_0 & (a_{22} + a_{33})\omega_0 & \left(a_{33} - \frac{a_{23}a_{31}}{a_{21}}\right)^2 + a_{23}\omega_0 \\ 0 & \omega_0 & a_{33} - \frac{a_{23}a_{31}}{a_{21}} \\ 0 & 0 & 1 \end{pmatrix}.$$

Denoting $\bar{\Pi}_1 = (F_4F_2)\Lambda_1(F_4F_2)^{\tau}$, $\bar{\Sigma}_1 = (F_4F_2)\Sigma_1(F_4F_2)^{\tau}$, (A.2) can then be equivalently transformed into the following equation: $\bar{\Pi}_1 + \bar{A}\bar{\Sigma}_1 + \bar{\Sigma}_1\bar{A}^{\tau} = 0$.

Similarly, by direct calculation, one obtains that $\widehat{\Pi}_1 = diag((a_{21}\omega_0)^2, 0, 0)$, and

$$\widehat{A} = \begin{pmatrix} -r_1 & -r_2 & -r_3 \\ 1 & 0 & 0 \\ 0 & 1 & 0 \end{pmatrix}, \quad \widehat{\Sigma}_1 = (a_{21}\omega_0)^2 \begin{pmatrix} \frac{r_2}{2(r_1r_2 - r_3)} & 0 & -\frac{1}{2(r_1r_2 - r_3)} \\ 0 & \frac{1}{2(r_1r_2 - r_3)} & 0 \\ -\frac{1}{2(r_1r_2 - r_3)} & 0 & \frac{r_1}{2r_3(r_1r_2 - r_3)} \end{pmatrix},$$

where r_1, r_2, r_3 are the same as those in Lemma 2.6. Considering

$$\bar{\Sigma}_1 = (a_{21}\omega_0)^2 \begin{pmatrix} \frac{1}{2r_1} & 0 & 0\\ 0 & 0 & 0\\ 0 & 0 & 0 \end{pmatrix} + (a_{21}\omega_0)^2 \begin{pmatrix} \frac{r_1}{2r_3(r_1r_2-r_3)} & 0 & -\frac{1}{2(r_1r_2-r_3)}\\ 0 & \frac{1}{2(r_1r_2-r_3)} & 0\\ -\frac{1}{2(r_1r_2-r_3)} & 0 & \frac{r_1}{2r_3(r_1r_2-r_3)} \end{pmatrix} := L_3 + L_4$$

(A.9)

(A.10)

one can therefore derive, by a similar method as that described in (A.7),

$$\begin{split} \Sigma_1 &= (F_4F_2)^{-1} \tilde{\Sigma}_1 [(F_4F_2)^{-1}]^{\tau} = (F_4F_2)^{-1} (L_3 + L_4) [(F_4F_2)^{-1}]^{\tau} \\ &= (F_4F_2)^{-1} L_3 [(F_4F_2)^{-1}]^{\tau} + (F_4F_2)^{-1} L_4 [(F_4F_2)^{-1}]^{\tau} \\ &= \begin{pmatrix} \frac{1}{2r_1} & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{pmatrix} + (F_4F_2)^{-1} L_4 [(F_4F_2)^{-1}]^{\tau} \\ &\coloneqq \Delta_{13} + (F_4F_2)^{-1} L_4 [(F_4F_2)^{-1}]^{\tau}. \end{split}$$

Noting that Δ_{13} , L_4 and $(F_4F_2)^{-1}L_4[(F_4F_2)^{-1}]^{\tau}$ are all positive semi-definite, one then has $\Sigma_1 \succeq \Delta_{13}$.

Consequently, by (A.4), (A.8) and (A.10), a constant $\eta_1 > 0$ always exists such that

$$\Sigma_1 \succeq \begin{pmatrix} \eta_1 & 0 & 0\\ 0 & 0 & 0\\ 0 & 0 & 0 \end{pmatrix}.$$
(A.11)

In addition, for the following two algebraic equations,

(i).
$$\Pi_2 + A\Sigma_2 + \Sigma_2 A^{\tau} = 0$$
, (ii). $\Pi_3 + A\Sigma_3 + \Sigma_3 A^{\tau} = 0$,
and letting $\widetilde{\Pi}_2 = F_5 \Pi_2 F_5^{\tau}$, $\widetilde{\Pi}_3 = F_6 \Pi_3 F_6^{\tau}$, $\widetilde{\Sigma}_2 = F_5 \Sigma_2 F_5^{\tau}$, $\widetilde{\Sigma}_3 = F_6 \Sigma_3 F_6^{\tau}$, $and A_2 = F_5 A F_5^{-1}$, $A_3 = F_6 A F_6^{-1}$, where

$$F_5 = \begin{pmatrix} 0 & 1 & 0 \\ 0 & 0 & 1 \\ 1 & 0 & 0 \end{pmatrix}, \quad F_6 = \begin{pmatrix} 0 & 0 & 1 \\ 1 & 0 & 0 \\ 0 & 1 & 0 \end{pmatrix}.$$

Noting that $\widetilde{\Pi}_2 = \widetilde{\Pi}_3 = \Pi_1$, by a method similar to that shown in (A.2), one can see that

$$\widetilde{\Sigma}_2 \succeq \begin{pmatrix} \eta_2 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{pmatrix}, \quad \widetilde{\Sigma}_3 \succeq \begin{pmatrix} \eta_3 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & 0 \end{pmatrix},$$

where the constants $\eta_2 > 0$ and $\eta_3 > 0$; that is to say,

$$\Sigma_{2} = F_{5}^{-1} \widetilde{\Sigma}_{2} F_{5}^{-1} \succeq \begin{pmatrix} 0 & 0 & 0 \\ 0 & \eta_{2} & 0 \\ 0 & 0 & 0 \end{pmatrix}, \quad \Sigma_{3} = F_{6}^{-1} \widetilde{\Sigma}_{3} F_{6}^{-1} \succeq \begin{pmatrix} 0 & 0 & 0 \\ 0 & 0 & 0 \\ 0 & 0 & \eta_{3} \end{pmatrix}.$$
(A.12)

In summary, it can be derived that

$$\Sigma_0 = \alpha_1^2 \Sigma_1 + \alpha_2^2 \Sigma_2 + \alpha_3^2 \Sigma_3 \succeq \begin{pmatrix} \alpha_1^2 \eta_1 & 0 & 0 \\ 0 & \alpha_2^2 \eta_2 & 0 \\ 0 & 0 & \alpha_3^2 \eta_3 \end{pmatrix}.$$

Given the above definitions and discussions, Σ_0 is a positive-definite matrix. This completes the proof.

(III). (SED Preliminaries): For the above complete probability space $\{\Omega, \mathscr{F}, \{\mathscr{F}_t\}_{t>0}, \mathbb{P}\}$, it is assumed that B(t) is an *n*-dimensional standard Brownian motion defined on it. Consider the following *n*-dimensional SDE,

$$dX(t) = f(X(t), t)dt + g(X(t), t)dB(t), \quad \text{for } t \ge t_0$$

, with the initial value $X(t_0) = X_0 \in \mathbb{R}^n$. A common differential operator \mathscr{L} is given by

$$\mathscr{L} = \frac{\partial}{\partial t} + \sum_{k=1}^{n} f_k(X, t) \frac{\partial}{\partial X_k} + \frac{1}{2} \sum_{i,j=1}^{n} \left[g^{\tau}(X, t) g(X, t) \right]_{ij} \frac{\partial^2}{\partial X_i \partial X_j}.$$

Letting the operator \mathscr{L} act on a function $V \in C^{2,1}(\mathbb{R}^n \times [t_0, +\infty]; \mathbb{R}^1_+)$, one has

$$\mathscr{L}V(X(t),t) = V_t(X(t),t) + V_x(X(t),t)f(X(t),t) + \frac{1}{2}trace[g^{\tau}(X(t),t)V_{xx}(X(t),t)g(X(t),t)],$$

where $V_t = \frac{\partial V}{\partial t}$, $V_x = (\frac{\partial V}{\partial x_1}, ..., \frac{\partial V}{\partial x_n})$ and $V_{xx} = (\frac{\partial^2 V}{\partial x_i \partial x_i})_{n \times n}$. If $X(t) \in \mathbb{R}^n$, one has

$$dV(X(t),t) = \mathscr{L}V(X(t),t)dt + V_X(X(t),t)g(X(t),t)dB(t).$$

References

- [1] Kermack WO, McKendrick AGM. A contribution to the mathematical theory of epidemics. Proc R Soc Lond A 1927;115:700-21.
- [2] Ma X, Wang W. A discrete model of avian influenza with seasonal reproduction and transmission. [Biol Dynam 2010;4:296-314.
- [3] Tchuenche JM, Nwagwo A, Levins R. Global behaviour of an SIR epidemic model with time delay. Math Methods Appl Sci 2007;30:733-49.
- [4] Hove-Musekwa SD, Nyabadza F. The dynamics of an HIV/AIDS model with screened disease carriers. Comput Math Method M 2015;10(4):287-305.
- [5] Lee H, Lao A, et al. Transmission dynamics and control strategies assessment of avian influenza a (H5N6) in the philippines. Infect Dis Model 2018;3:35-59. [6] Tuncer N, Martcheva M. Modeling seasonality in avian influenza H5N1. J Biol Syst 2013;21(4):1-30.
- [7] Cai L, Wu J. Analysis of an HIV/AIDS treatment model with a nonlinear incidence. Chaos Solitons Fractals 2009;41(1):175-82.
- [9] Liu X, Takeuchi Y, Iwami S. SVIR epidemic models with vaccination strategies. J Theore Bio 2008;253:1–11.

[10] Li J, Ma Z. Qualitative analyses of SIS epidemic model with vaccination and varying total population size. Math Comput Model 2002;35:1235-43.

- [11] Gao S, Ouyang H, Nieto J. Mixed vaccination strategy in SIRS epidmeic model with seasonal variability on infection. Int J Biomath 2011;4:473-91.
- [12] Chen F. A susceptible-infected epidemic model with voluntary vaccinations. | Math Biol 2006;53:253-72.
- [13] Al-Darabsah I. Threshold dynamics of a time-delayed epidemic model for continuous imperfect-vaccine with a generalized nonmonotone incidence rate. Nonlinear Dyn 2020;101:1281-300.
- [14] Anderson RM, May RM. Infectious diseases in humans: dynamics and control. Oxford: Oxford University Press; 1991.
- [15] Hethcote HW. The mathematics of infectious diseases. SIAM Rev 2000;42:599-653.
- [16] Yang Q, Jiang D, Shi N, Ji C. The ergodicity and extinction of stochastically perturbed SIR and SEIR epidemic models with saturated incidence. J Math Anal Appl 2012:388:248-71.
- [17] Wang Y, Liu J, Liu L. Viral dynamics of an HIV model with latent infection incorporating antiretroviral therapy. Adv Differ Equ 2016;225:1-15.
- [18] Liu S, Pang L, Ruan S, Zhang X. Global dynamics of avian influenza epidemic models with psychological effect. Comput Math Method Med 2015;2015:913726.
 [19] Xiao D, Ruan S. Global analysis of an epidemic model with nonmonotone incidence rate. Math Biosci 2007;208(2):419–29.
- [20] Anderson R, May R. Infectious diseases of human: dynamics and control. Oxford: Oxford University Press; 1991.
- [21] Anderson R, May R. Population biology of infectious diseases. Berlin, Heidelberg New York:: Springer-Verlag; 1982.
- [22] Ma Z, Zhou Y, et al. Modeling and dynamic of infectious disease. Beijing: Higher Education Press; 2009. (In Chinese)
- [23] Truscott JE, Gilligan CA. Response of a deterministic epidemiological system to a stochastically varying environment. Proc Natl Acad Sci 2003;100:9067-72.
- [24] Liu Z. Dynamics of positive solutions to SIR and SEIR epidemic models with saturated incidence rates. Nonlin Anal Real World Appl 2013;14(3):1286-99.
- [25] Zhu P, Wang X, Li S, Guo Y, Wang Z. Investigation of epidemic spreading process on multiplex networks by incorporating fatal properties. Appl Math Comput 2019-359-512-24
- [26] Jia D, Wang X, Song Z, Rom0ć, Li X, Jusup M, Wang Z. Evolutionary dynamics drives role specialization in a community of players. J R Soc Interface 2020;17:168. 20200174
- [27] Mao X, Marion G, Renshaw E. Environmental Brownian noise suppresses explosions in population dynamics. Stoch Pro Appl 2002;97(1):95–110.
- [28] Zhou B, Zhang X, Jiang D. Dynamics and density function analysis of a stochastic SVI epidemic model with half saturated incidence rate. Chaos Soliton Fractals 2020.137.109865
- [29] Qi K, Jiang D. The impact of virus carrier screening and actively seeking treatment on dynamical behavior of a stochastic HIV/AIDS infection model. Appl Math Model 2020:85:378-404.
- [30] Khan T, Khan A. The extinction and persistence of the stochastic hepatitis b epidemic model. Chaos Soliton Fractals 2018;108:123-8.
- [31] Wang Y, Jiang D. Stationary distribution of an HIV model with general nonlinear incidence rate and stochastic perturbations. J Frank Inst 2019;356:6610–37.
- [32] Caraballo T, Fatini ME, Khalifi ME. Analysis of a stochastic distributed delay epidemic model with relapse and gamma distribution kernel. Chaos Soliton Fractals 2020:133:109643.
- [33] Cai Y, Kang Y. A stochastic epidemic model incorporating media coverage. Commun Math Sci 2015;14:893–910.
- [34] Shi Z, Zhang X, Jiang D. Dynamics of an avian influenza model with half-saturated incidence. Appl Math Comput 2019;355:399-416.
- [35] Yang Q, Jiang D, Shi N, Ji C. The ergodicity and extinction of stochastically perturbed SIR and SEIR epidemic models with saturated incidence. J Math Anal Appl 2012;388:248-71.
- [36] Cai Y, Kang Y. A stochastic epidemic model incorporating media coverage. Commun Math Sci 2015;14:893–910.
- [37] Zhao Y, Jiang D. The threshold of a stochastic SIS epidemic model with vaccination. Appl Math Comput 2014;243:718-27.
- [38] Liu Q, Jiang D, Hayat T, Alsaedi A. Threshold behavior in a stochastic delayed SIS epidemic model with vaccination and double diseases. J Frank Inst 2019;356:7466-85.
- [39] Zhang X, Jiang D. Dynamics of a stochastic SIS model with double epidemic diseases driven by Lévy jumps. Physica A 2017;471:767-77.
- [40] Zhang X, Jiang D. Dynamical behavior of a stochastic SVIR epidemic model with vaccination. Physica A 2017;483:94-108.
- [41] Mao X. Stochastic differential equations and applications. Chichester: Horwood Publishing; 1997
- [42] Has'miniskii RZ. Stochastic stability of differential equations. The Netherlands;: Sijthoff Noordhoff, Alphen aan den Rijn; 1980.
- [43] Ma Z, Zhou Y, Li C. Qualitative and stability methods for ordinary differential equations. Science Press; 2015.
- [44] Gardiner CW. Handbook of stochastic methods for physics. Chemistry and the natural sciences. Springer Berlin; 1983.
- [45] Roozen H. An asymptotic solution to a two-dimensional exit problem arising in population dynamics. SIAM J Appl Math 1989;49:1793.
- [46] Higham DJ. An algorithmic introduction to numerical simulation of stochastic differential equations. SIAM Rev 2001;43:525-46.
- [47] Arino J, Mccluskey CC, Driessche P. Global results for an epidemic model with vaccination that exhibits backward bifurcation. SIAM J Appl Math 2003;64(1):260276.
- [48] Liu Q, Jiang D, Shi N, Hayat T, Alsaedi A. Asymptotic behavior of stochastic multi-group epidemic models with distributed delays. Physica A 2017;467:527-41.
- [49] Li X, Gray A, Jiang D, Mao X. Sufficient and necessary conditions of stochastic permanence and extinction for stochastic logistic populations under regime switching. J Math Anal Appl 2011;376:11-28.
- [50] Zhang X, Jiang D, Alsaedi A. Stationary distribution of stochastic SIS epidemic model with vaccination under regime switching. Appl Math Lett 2016;59:87-93.